Appendix A

TP-201.2

DETERMINATION OF EFFICIENCY OF PHASE II VAPOR RECOVERY SYSTEMS OF DISPENSING FACILITIES

California Environmental Protection Agency



Air Resources Board

Vapor Recovery Test Procedure

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Adopted: April 12, 1996

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1 APPLICABILITY

A set of definitions common to all certification and test procedures is in:

D-200 Definitions for

Certification Procedures and

Test Procedures for

Vapor Recovery Systems

For the purpose of this procedure, the term "ARB" refers to the State of California Air Resources Board, and the term "ARB Executive Officer" refers to the Executive Officer of the ARB or his or her authorized representative or designate.

1.1 General

This procedure applies to the determination of Phase II vapor recovery system efficiency at dispensing facilities by mass balance principles. This procedure applies to any vapor emissions associated with the dispensing of any fluid, although it is written to reflect application to the hydrocarbon (HC) vapors associated with the dispensing of gasoline.

1.2 Modifications

Modification of this procedure may be necessary for vapors and fluids other than the hydrocarbon vapors associated with the dispensing of gasoline.

Any modification of this method shall be subject to approval by the ARB Executive Officer.

2 PRINCIPLE AND SUMMARY OF TEST PROCEDURE

The purpose of this test procedure is to determine the percent vapor recovery efficiency for a vapor recovery system at a dispensing facility. The percent vapor

recovery efficiency is the percent of vapors displaced by dispensing which are recovered by a vapor recovery system rather than emitted to the atmosphere.

2.1 Principle

This is done by simultaneously measuring the vapor mass flux through four significant areas:

- $m_{(1)}$ = the mass flux through openings at the dispensing interface,
- $m_{(2)}$ = the mass flux through the vapor return line,
- $m_{(3)}$ = the mass flux through the dispensing facility vent, and
- $m_{(4)}$ = the mass flux through the assist processor.

The percent vapor recovery efficiency is determined as follows:

% vapor recovery efficiency =
$$\frac{m_{(2)} - [m_{(3)} + m_{(4)}]}{[m_{(2)} + m_{(1)}]} \times 100\%$$

2.2 Summary

As required to determine an emissions parameter and except where otherwise specified, the equipment and procedures specified in the following test methods shall be used.

EPA Method 2A	Direct Measurement of Gas Volume Through Pipes and Small Ducts
EPA Method 2B	Determination of Exhaust Gas Volume Flow Rate From Gasoline Vapor Incinerators
EPA Method 18	Measurements of Gaseous Organic Compound Emissions by Gas Chromatography
EPA Method 25A	Determination of Total Gaseous Organic Compound Emissions Using a Flame Ionization Detector
EPA Method 25B	Determination of Total Gaseous Organic Compound Emissions Using a Nondispersive Infrared Analyzer

3 BIASÉS AND INTERFERENCES

3.1 Vehicle Biases and Interferences

3.1.1 Inclusion of Vehicles in Test Procedure

A representative vehicle matrix shall be determined for the subject facility according to TP-201.28.A

3.1.2 Exclusion of Certain Vehicle Results from Test Results

Include the following vehicles in this test procedure, but exclude results for such vehicles from any determinations of compliance with or violation of the certification criterion. Report results for such vehicles separately, with a description of the likely causes for their failing to meet any requirements specified below.

For the purpose of determinations of compliance with or violation of the certification criterion, exclude vehicles with:

(1) non-conformance with other applicable requirements,

results for vehicles with fillpipe access zones which do not conform with applicable specifications and requirements (vehicles shall arrive at the facility with properly positioned fillpipe caps and leaded fuel restrictors appropriate for each vehicle);

(2) non-conformance with vehicle leak check requirement,

Note: This vehicle leak check requirement may be waived, on a vehicle-by-vehicle basis, upon determination by the ARB Executive Officer that the vehicle matrix required by TP-201.2A can not otherwise be filled.

This vehicle leak check requirement may be waived, on a system-by-system basis, upon determination by the ARB Executive Officer that the system is expected to always maintain negative gauge pressure in all vehicle tanks during all dispensing episodes.

results for vehicles which do not pass the vehicle leak check requirement (>0.01 cfm); and

(3) non-conformance with sleeve leak check requirement, and

results for vehicles with leak detector readings (per EPA Method 21) above 0.1% LEL within one inch (2.5 cm) outside the sampling sleeve; and

(4) inadequate dispensed volume.

results for vehicles into which less than four gallons are dispensed.

3.2 Facility Biases and Interferences

3.2.1 Static Pressure Performance

The subject dispensing facility shall demonstrate compliance with the appropriate static pressure performance standard as required by CP-201.

3.2.2 Representative Facility Operating Matrix

The subject facility shall operate within the matrix of conditions established in the specifications of certification. See CP-201 § 3 and § 5.

During certification testing, any conditions of installation, operation, and maintenance which deviate from such specifications, shall be recorded and included as amendments to the specifications of certification. Subsequent to such certification, any conditions which occur outside such specifications (for any facility installed, operating, and maintained on the basis of such certification) shall constitute a violation of the specifications of certification.

3.2.3 Dispensed Liquid Characteristics

Some unusual situations may require a more extended calibration protocol if, in a certain case, the speciation of vapors is significantly different than that for gasoline vapors. Two alternative approaches, both of which must be demonstrated in practice and approved as modifications to this procedure are suggested:

- (1) calibrate all analyzers to appropriate dilutions of a Tedlar® bag sample taken from the headspace of the facility involved in the proposed test, or
- (2) use the same make and model of analyzer at each test point while minimizing the amount of vapor taken from the vapor return line. At the vapor return line test point, this may require a high flow (to reduce time lag) sample extraction and return loop and a low flow (to satisfy analyzer requirements) sample line from the sample loop to the analyzer.

3.3 Equipment Biases and Interferences

Alternatives to the required equipment can be as good or better in certain testing circumstances. Such alternatives shall only be used subject to prior written approval by the ARB Executive Officer, as required in § 13.

A primary example of such an alternative is the use of NDIR instead of FID at Test Point 1 for analyzing the sleeve sample. There NDIR can be used based on

data and documentation which show, to the satisfaction of the ARB Executive Officer, satisfactory control of biases and interferences due to the use of the equipment.

4 SENSITIVITY, RANGE, AND PRECISION

The measurements of concentration and volumetric parameters required by this test procedure are well within the limits of sensitivity, range, and precision of the specified equipment.

5 EQUIPMENT

Equipment specifications are given below and some equipment configurations are shown in Figures 1 through 21.

5 (1) Hydrocarbon Analyzer

The default mode of determining hydrocarbon concentrations in this procedure is a determination of total hydrocarbon concentration as propane. Alternative test procedures for determining non-methane hydrocarbon concentration have been validated for some applications and may be used, subject to the approval of the ARB Executive Officer. Such procedures typically determine both the total hydrocarbon concentration and methane concentration; non-methane hydrocarbon concentration is then determined as the difference.

The range of any hydrocarbon analyzer shall be selected such that the maximum concentration measured is no more than 90 percent of the range and the average concentration is no less than 10 percent of the range.

Any sampling and analysis system using a non-dispersive infrared detector (NDIR) shall be designed so that 100% of the sample that is extracted for analysis can be returned, unaltered, to the sample manifold.

An analyzer with a NDIR with selected filters to block methane measurement shall be used when the efficiency is to be calculated for non-methane hydrocarbon and when the system under test is small such that extracting a sample for a FID analyzer will affect the system operating parameters. When using a NDIR instrument for total hydrocarbon measurements, a second channel must be present to measure the methane concentration or the instrument filters must be such that total hydrocarbon is measured.

Any sampling and analysis system using a flame ionization detector (FID) can not be designed so that 100% of the sample that is extracted for analysis can be returned, unaltered, to the sample manifold, because the operation of the FID significantly alters the portion of the sample which is analyzed.

An analyzer with a FID may be used for the test when a measurement is for total hydrocarbon and there is no requirement for returning sample, unaltered, to the sample manifold. An important example is the total hydrocarbon measurement on the diluted sample from a test sleeve which has captured

transfer emissions from the nozzle fillpipe interface. In this case, the transfer emissions are on their way to the atmosphere normally, so there is no need to return them to a sample manifold.

5 (2) Carbon Monoxide Analyzer

Use a NDIR analyzer for measurement of exhaust CO concentrations. To the extent practical, the analyzer range shall be selected such that the maximum concentration measured is no more than 90 percent of the range and the average concentration is no less than 10 percent of the range.

5 (3) Carbon Dioxide Analyzer

Use a NDIR analyzer for measurement of exhaust CO_2 concentrations. The analyzer range shall be selected such that the maximum concentration measured is no more than 90 percent of the range and the average concentration is no less than 10 percent of the range.

5 (4) Volume

Use a calibrated positive displacement gas volume meter or a turbine meter for measurement of volumetric flow rate.

Use rotary type positive displacement meter(s) with a back pressure limit (BPL) less than:

- 1.10 inches water column at a flowrate of 3,000 CFH down to 0.05 inches water column at a flowrate of 30 CFH for a meter with a rating over 1000 CFH and
- 0.70 inches water column at a flowrate of 800 CFH down to 0.04 inches water column at a flowrate of 16 CFH for a meter with a rating of or under 1000 CFH.

Meter(s) shall be equipped with taps accommodating the following equipment:

- (1) taps on the inlet side for
 - (a) a thermocouple with a range of 0 to 150 °F and
 - (b) a pressure gauge with a range providing absolute pressure readings within 10 to 90% of the range (more than one gauge shall be used, if necessary) and
- (2) taps on the inlet and outlet sides for a differential pressure gauge with a range of 0 to < 2x BPL (i.e. full scale shall be less than twice the back pressure limit) or any other range appropriate to allow detection of a pressure drop greater than the BPL.

5 (5) Pressure

Use a pressure measuring device (transducer, inclined manometer or Magnahelic gauge) with a design range suitable for the pressure being measured. The tap for the pressure measurement shall be located on the sample coupling attached to the inlet of the volume meter.

5 (6) Temperature

Use a temperature measuring device (thermocouple or mercury in glass thermometer) with a design range suitable for the temperature being measured. The tap for the temperature measurement shall be located on the sample coupling attached to the inlet of the volume meter.

5.1 Equipment for Test Point 1 (Nozzle Sleeve)

5.1.1 Vehicle Leak Check Equipment

The following equipment is necessary to perform required vehicle leak checks; or to demonstrate that alternative equipment will perform equivalently (see "ALTERNATIVE TEST PROCEDURES" section).

5.1.1.1 Fillpipe Interface

A fillpipe interface shall be used which provides a seal at the fillpipe outlet except for:

- (1) tubing for pressurizing the fillpipe and vehicle tank with nitrogen and
- (2) tubing for connection to a pressure transducer which can register the pressure in the fillpipe and vehicle tank.

5.1.1.2 Flowmeter and Pressure Transducer

A flowmeter and pressure transducer shall be used which are appropriately sized for measuring 0.01 cfm and one-half (0.5) inches water (gauge) at the fillpipe interface.

5.1.1.3 Pressure System

The pressure system for the vehicle leak check shall consist of a nitrogen bottle (2000 psig, commercial grade), a control valve for regulating the bottle pressure to 1 psig, a needle valve, two Magnahelic gauges (0 - 30 and 0 - 10 inches water) for determining the pressure upstream and downstream of the needle valve, and a hose for supplying pressure to the vehicle tank. The pressure system shall provide for monitoring the pressure in the fillpipe and vehicle tank during the vehicle leak check.

5.1.2 Sleeve Leak Check Equipment

A volatile organic compound detector which complies with the requirements of EPA Method 21 shall be used.

5.1.3 Implements

5.1.3.1 Sleeve

The sleeve is designed for vapor sampling at the dispensing area. An example design for the sleeve is shown in Figures 6 through 8.

Other designs may be used which accommodate different dispensing area geometries, subject to the requirement that other designs yield no more pressure drop at five cubic feet per minute (cfm) air flow than the design shown. Compliance with this requirement must be documented in the test report required in the section, "REPORTING RESULTS".

The design shown has been tested, at 5 cfm with -0.005 "WC gauge pressure inside the sleeve, during use in a balance nozzle application. The comparison standard may differ in other dispensing geometries.

5.1.3.2 Sleeve Sample Tubing

The sample tube connecting the sleeve to its instrumentation shall be as flexible and lightweight as practical so that the behavior of the nozzle operator is minimally affected by testing activities. It is not necessary to return the unanalyzed portion of sample flow back to the GDF vapor recovery system.

In general, only a portion of the sleeve flow is used for analysis. Most analyzers sample at a flow rate far below the 5 cfm sleeve flow rate. In such a case, sleeve sample tubing must be configured so that a portion of the sleeve flow is representatively sampled by the analyzer at conditions suitable for the analyzer requirements.

5.1.3.3 Sleeve Sample Pump

Note: The sample flow rate must always be high enough to prevent the sleeve leak check from registering more than 0.1% LEL (2,100 ppm as propane).

Use a carbon vane (or equivalently non-contaminating) pump to minimize contamination of the sample.

The pump must be capable of pulling about 5 cfm, but lower flow rates are acceptable subject to the following requirement:

The pressure drop is typically a few inches Hg, depending on tubing and fittings.

5.1.4 Instruments

5.1.4.1 Hydrocarbon Concentration

Use an FID with full scale values of 1.00% and 10.0%. Perform span and calibration checks with propane standards.

5.1.4.2 Volume (See § 5.4)

5.1.4.3 Pressure

Use a pressure measuring device (transducer, inclined manometer or Magnahelic gauge) with a design range suitable for the pressure being measured. The tap for the pressure measurement shall be located on the sample coupling attached to the inlet of the volume meter.

5.1.4.4 Temperature

A transducer with an initial design range of 0 - 150°F in a thermocouple design, depending on the sleeve tubing chosen. The tap must be near the HC instrument inlet.

5.2 Equipment for Test Point 2 (Vapor Return)

5.2.1 Implements

5.2.1.1 In-line Plumbing

Design goals for plumbing arrangements, regardless of GDF, are:

- (1) practically minimize length of vapor return line between the nozzle and the sample point for Test Point 2; do this to minimize problems related to entry of condensation from the vapor return line into the Test Point 2 sample line;
- (2) practically minimize pressure drop across in-line plumbing; and
- (3) return any unanalyzed sample to the GDF vapor return line.

5.2.1.2 Fittings

Plumbing shall be designed for easy adaptability to co-axial, twin hose, and any other GDF configurations which may be encountered. A one inch (i.d.) ball valve shall isolate the vapor return line from other implements.

5.2.2 Instruments

5.2.2.1 Hydrocarbon Concentration

Use a NDIR with a full scale value of 100.0%, or a lower value which is known to be above the maximum concentration possible at test

conditions. Perform span and calibration checks with appropriate propane standards.

5.2.2.2 Volume (See § 5.4)

5.2.2.3 Pressure

Use a pressure measuring device (transducer, inclined manometer or Magnahelic gauge) with a design range suitable for the pressure being measured. The tap for the pressure measurement shall be located on the sample coupling attached to the inlet of the volume meter.

5.2.2.4 Temperature

Use a transducer with an initial design range of 0 - 150°F in a thermocouple design.

5.3 Equipment for Test Point 3 (Vent)

5.3.1 Implements

Unanalyzed sample shall be returned to the system to avoid perturbation of the system pressure.

5.3.2 Instruments

5.3.2.1 Hydrocarbon Concentration

Upstream from any final point of release to the atmosphere, use a NDIR with a full scale value of 100.0%, or a lower value which is known to be above the maximum concentration possible at test conditions. Perform span and calibration checks with appropriate propane standards.

Sample may be collected downstream of the normal point of release to the atmosphere in situations where upstream techniques are precluded. Some special sealed sleeve or other means of gathering and delivering sample to the analyzer must be employed; before testing commences, any such sleeve or means must pass the leak check criteria given in the procedures provided for leak checking the nozzle sleeve. Tape and polyethylene bags have been used to successfully meet this requirement.

5.3.2.2 Volume (See § 5.4)

5.3.2.3 Pressure

Use a pressure measuring device (transducer, inclined manometer or Magnahelic gauge) with a design range suitable for the pressure being measured. The tap for the pressure measurement shall be located on the sample coupling attached to the inlet of the volume meter.

5.3.2.4 Temperature

Use a transducer with an initial design range of 0 - 150°F in a thermocouple design.

5.4 Equipment for Test Point 4 (Assist Processor Exhaust)

See section, "ALTERNATIVE TEST PROCEDURES" if equipment specified above is not applicable.

6 CALIBRATION PROCEDURE

A record of all calibrations shall be maintained.

6.1 Analyzers

Follow the manufacturer's instructions concerning warm-up time and adjustments. On each test day prior to testing, zero the analyzer with a zero gas and span with a known concentration of calibration gas at a level near the highest concentration expected. Perform an intermediate zero and span calibration approximately 2 hours after the initial calibration and at any time a calibration drift is evident. Check for zero and span calibration drift at the end of the test period. All calibrations and adjustments shall be documented.

6.2 Volume Meters

Meters shall be calibrated on an annual basis.

6.3 Pressure Transducers

Calibrate pressure transducers prior to testing and immediately following the test period with a static pressure calibrator for a range of -3 to +3 inches water or appropriate range of operation; or calibrate pressure transducers in accordance with manufacturer's specifications and provide a copy of such specifications in the Certification Test Report.

6.4 Temperature Transducers

Calibrate temperature transducers every six months using ambient air, the temperature of which is determined by a NIST traceable mercury-glass thermometer.

7 PRE-TEST PROTOCOL

7.1 Location of Test Site

Prototype systems will be located within 100 miles of Sacramento for testing. Other locations may be accepted at the discretion of the ARB Executive Officer.

7.2 Specification of Test, Challenge, and Failure Modes

The specification of test, challenge, and failure modes such as the number of liquid transfer episodes, volume and volumetric rate of liquid transfer, storage tank volumes, etc. shall be done according to the principles of CP-201 § 5 for the testing and evaluation of vapor recovery equipment.

7.3 System and Facility Preparation

The required preliminary evaluation shall set the final requirements for facility preparation. The dominant principle shall be that testing activities minimally alter facility and system conditions. As the installation of test equipment can alter facility and system values for critical parameters, the following final preparation procedures shall be applied, subject to determination by the ARB Executive Officer of more effective alternatives for some procedures and the only practical alternatives for other procedures:

(1) Install all equipment and wait at least 16 hours before testing. Until then, provide conditions which minimally disturb facility and system operations due to the presence of such equipment for such time; or

Warning: the following alternative shall only be used after a determination, per the preliminary evaluation, that system pressure is the only system parameter disturbed by equipment installation and that volumetric flow from the system can be monitored by procedures which minimally disturb facility and system operations.

(2) install all equipment and wait until a determination of a flow of 0.1 ACF from the system before testing. Until then, provide conditions which minimally disturb facility and system operations due to the presence of such equipment for such time.

7.4 Testing Sequence

Note:

For the applicant, this test procedure is the most costly required test procedure. As the applicant's candidate system must pass all required tests, costs for the applicant can be minimized by performing all other procedures before the "100 car" test specified in this procedure and TP-201.2A. In this way, if the system fails one of the less costly tests, required redesign and refabrication of the system can proceed at the least cost.

The testing sequence shall be as follows:

7.4.1 Other Required Test Procedures

All other required test procedures shall be performed before the application of TP-201.2A for the 100 car test specified in this procedure.

7.4.2 Other Required Aspects of this Test Procedure

All other required aspects of this test procedure shall be performed before the application of TP-201.2A for the 100 car test specified in this procedure, e.g. testing idle nozzle episodes.

8 TEST PROCEDURE

The facility and system shall be prepared to operate according to any specified test, challenge, and failure modes.

In this section, the term "vent" and the specified procedures for testing vents shall also apply to any assist processor with which such procedures are compatible. Procedures are also specified for incinerator type assist processors. Any assist processor which is incompatible with the application of these procedures shall not be certified until the compatibility requirements of the certification procedures are met.

8.1 Test Locations

Figure 1 illustrates mass flux test locations.

8.1.1 Test Point 1 (Nozzle Sleeve)

Figure 2 emphasizes the mass flux test location for Test Point 1 (Nozzle Sleeve).

8.1.1.1 Vehicle Leak Check Procedure

Three different procedures are acceptable for checking leaks in vehicle tanks and plumbing.

(1) Nitrogen Pressurization

Figure 3 illustrates the following procedure, which is necessary to perform required vehicle leak checks; or to demonstrate that an alternative procedure will perform equivalently (see "ALTERNATIVE TEST PROCEDURES" section).

- (a) Connect equipment for vehicle leak check to vehicle fillpipe.
- (b) Open main valve on the nitrogen supply bottle and adjust the needle valve until the pressure in the fillpipe reaches one-half (0.5) inches water (gauge). If such pressure can not be maintained for 15 seconds, record an unacceptable vehicle leak for the subject vehicle.
- (c) Determine the leak rate by either timing a volume of 0.1 ft³ or by observing for 15 seconds, whichever results in a smaller volume being transferred to the vehicle tank.

 Record readings.

- (d) Disconnect equipment from the vehicle fillpipe and proceed with further test procedures.
- (e) If a leak-rate greater than 0.01 cfm, record an unacceptable vehicle leak for the subject vehicle.

(2) Manual Compression

Figure 4 illustrates the following procedure.

- (a) Use a vapor tight, sealed, compressible device with an attached pressure gauge and seal the device against the vehicle fill pipe interface.
- (b) Compress the device in a repeatable and controlled manner and record readings from the pressure gauge.
- (c) Determine vehicle leak check status by comparing pressure readings with a calibration chart which must be developed independently, for each compression device and tester, as specified below:
 - (i) Perform the specified vehicle leak check procedure and the alternative compression procedure on a series of vehicles.
 - (ii) Correlate the readings from the specified vehicle leak check procedure and the alternative compression procedure which relate to passing and failing the vehicle leak check requirement. Include all significant variables in the correlation.
 - (iii) Construct a calibration chart from the correlations in (ii), indicating those readings from the alternative compression procedure which correlate with passing and failing the vehicle leak check requirement.

(3) Manual De-compression

Figure 5 illustrates the following procedure.

- (a) Prepare to listen for a sound of vapor de-compression from the vehicle tank and fillpipe, before removing a vehicle fillpipe cap.
- (b) Remove the cap in a quick, repeatable, and controlled manner and listen for a sound of vapor de-compression from the vehicle tank and fillpipe. Record a positive or negative reading of the occurrence of such sound when the cap is removed.

- (c) Determine vehicle leak check status by comparing positive readings with a calibration chart which must be developed independently, for each de-compression tester, as specified below:
 - (i) Perform the specified vehicle leak check procedure and the alternative de-compression procedure on a series of vehicles.
 - (ii) Correlate the readings from the specified vehicle leak check procedure and the alternative de-compression procedure which relate to passing and failing the vehicle leak check requirement. Include all significant variables in the correlation.
 - (iii) Construct a calibration chart from the correlations in (ii), indicating those readings from the alternative de-compression procedure which correlate with passing and failing the vehicle leak check requirement.

8.1.1.2 Nozzle Sleeve Assembly

The sleeve must be sampling around all potential vapor leak paths at all times during testing including:

dispensing periods; and

"idle nozzle" periods

as explained below, after a description of the sleeve leak check procedure.

(1) Sectional View of Sleeve

Figure 6 illustrates a sectional view of a nozzle sleeve.

(2) Axial View of Sleeve

Figure 7 illustrates an axial view of a nozzle sleeve.

(3) View of Sleeve on Nozzle

Figure 8 illustrates a view of a nozzle sleeve on a nozzle.

The sleeve must be sampling around all potential vapor leak paths at all times during testing including:

- (a) dispensing periods; and
- (b) "idle nozzle" periods

as explained below, after a description of the sleeve leak check procedure.

8.1.1.3 Leak Check of Sleeve

At least once during each dispensing period and once during "hang time" (as soon as practical after nozzle "hang-up"), readings must be taken with a leak detector per EPA Method 21. If possible, adjust the sleeve so that readings are below 0.1% LEL (2,100 ppm as propane) during within one inch (2.5 cm)-outside the sampling sleeve.

(1) View of Combustible Gas Detector

Figure 9 illustrates a view of a combustible gas detector.

(2) View of Combustible Gas Detector in Use

Figure 10 illustrates a view of a combustible gas detector in use.

8.1.1.4 Nozzie Sleeve Measurements

The sleeve temperature and pressure measurements must be taken from a sample manifold attached to the inlet of the volume meter on the sleeve sampling system. The hydrocarbon sample shall be taken at the exhaust side of the volume meter.

In the interest of reducing the amount of chart paper to be recorded and read and reducing the wear on pumps, and at the discretion of the ARB Executive Officer, the chart drive and pumps may be turned off whenever the sleeve sample concentration drops below 100 ppm. If this option is exercised by the test team, they must stand ready to resume operation of all sampling equipment immediately after the sample concentration climbs above 100 ppm. Also, the start and stop times for such periods during which the chart drive is off must be clearly marked on the chart record. See "ALTERNATIVE TEST PROCEDURES" section.

(1) Volume Measurement

Figure 11 illustrates equipment for volume measurements of samples from the nozzle sleeve.

(2) Concentration Measurement

Figure 12 illustrates equipment for concentration measurements of samples from the nozzle sleeve.

8.1.1.4.1 Dispensing Episodes

The tester begins data collection for a dispensing episode with the insertion of the nozzle into the vehicle and continues until the end of the "response time" which is defined per EPA Method 21 as follows:

Introduce zero gas into the sleeve until the analyzer reading has stabilized, then switch quickly to the specified calibration gas (§ 6.1). Measure the time interval from switching to attainment of 90% of the final stable analyzer reading. Perform this test sequence three times, calculate the average, and define the result as the "response time."

The nozzle user is to dispense normally and terminate dispensing in the user's customary manner. The tester shall also instruct the user that upon deciding that termination is complete, the nozzle user shall so declare for the tester to hear.

To achieve this, and prior to nozzle insertion and for every dispensing episode, the tester shall provide simple, clear instructions to the nozzle user. The instructions shall be the same for each nozzle user.

After hearing that the user has terminated dispensing, the tester waits for the response time and then ends data collection for the dispensing episode.

The sleeve must always be at the fillpipe/nozzle interface for sample collection during any dispensing episode.

Sample at a nominal flow rate of 5 cfm, or less subject to the requirement that the sleeve leak check is less than 0.1% LEL (2,100 ppm as propane).

8.1.1.4.2 Idle Nozzle Episodes

An idle nozzle episode is any time other than a dispensing episode.

In the interest of improving the accuracy of idle nozzle data and reducing the wear on pumps, and at the discretion of the ARB Executive Officer, idle nozzle data may be taken from vehicles other than those in the required vehicle matrix for efficiency testing. Accuracy can be improved and pump wear reduced by avoiding the need to frequently change sample pump speed (to change from one dilution for the higher concentration dispensing sample to another dilution for the lower concentration idle nozzle sample). This option shall only be allowed if an adequately representative sample of vehicles is used for idle nozzle sampling. See "ALTERNATIVE TEST PROCEDURES" section.

Because idle nozzle emissions do not involve the nozzle/fillpipe interface, the determination of a vehicle matrix per TP-201.2A is not necessary.

(1) Idle Nozzle Screening Test Procedure

- (a) Follow the instructions for the Leak Check of the Sleeve, but use the leak detector to determine if the nozzle is "leaking" idle nozzle emissions.
 - (i) Non-compliance is indicated by a reading at or above 2,100 ppm as propane (0.1% LEL).
 - (ii) Compliance is indicated when the procedures of EPA Method 21 have been followed and all readings are below 2,100 ppm as propane (0.1% LEL).
- (b) Collect such data for up to twenty-five (25) idle nozzle episodes.
 - (i) If a non-complying episode occurs, proceed to the Idle Nozzle Sleeve Test Procedure below.
 - (ii) If only complying episodes occur for the first twenty-five (25) idle nozzle episodes, record a determination of compliance with the idle nozzle test requirement; indicate in the Certification Report that idle nozzle emissions were not detected.

(2) Idle Nozzle Sieeve Test Procedure

- (a) Empirically determine, record, and employ a sample pump flow rate and analyzer range appropriate for any detectable idle nozzle emissions from the system.
- (b) Using the procedures and calculations for Test Point 1, prepare to collect data in order to calculate results in units of pounds of hydrocarbon emitted per thousand gallons of fuel dispensed.
- (3) Collect such data for twenty-five (25) idle nozzle episodes.
 - (a) If the overall result, in units of pounds of hydrocarbon emitted per thousand gallons of fuel dispensed, for all mass emitted divided by all fuel dispensed is over 0.42 pounds per thousand gallons (0.42 #/E3G), disapprove the application per CP-201 §2.2; the applicant must re-apply for

further consideration of the system except as (4) below is applied.

- (b) If the overall result is at or under 0.42 #/E3G, record the result for further calculations and proceed with the remainder of the required tests.
- (4) At the applicants option, collect such data for seventy-five (75) more idle nozzle episodes; follow the instructions provided in step (3) (a) to obtain the overall result for all one hundred (100) data sets.
 - (a) If the overall result, in units of pounds of hydrocarbon emitted per thousand gallons of fuel dispensed, for all mass emitted divided by all fuel dispensed is over 0.42 pounds per thousand gallons (0.42 #/E3G), disapprove the application per CP-201 §2.2; the applicant must re-apply for further consideration of the system.
 - (b) If the overall result is at or under 0.42 #/E3G, record the result for further calculations and proceed with the remainder of the required tests.

8.1.2 Test Point 2 (Vapor Return)

Figure 13 emphasizes mass flux test locations for Test Point 2 (Vapor Return Line).

The vapor return line sample and temperature and pressure measurements must be taken from a sample manifold attached to the inlet of the volume meter which has been inserted at a break in the vapor return line. The break is usually at the vapor hose connection to the vapor riser from under the pavement. When options are available, the sampling location shall be the shortest practical downstream distance from the nozzle to minimize vapor condensation upstream of the sampling location. Unaltered sample shall be returned to the sample manifold.

8.1.2.1 Volume Measurement, General

Figure 14 generally illustrates equipment for volume measurements of samples from the vapor return line.

8.1.2.2 Volume Measurement, Single Vapor Return Line

Figure 15 specifically illustrates metering equipment for volume measurements of samples from a single vapor return line.

8.1.2.3 Volume Measurement, Manifolded Vapor Return Lines

Figure 16 specifically illustrates metering equipment for volume measurements of samples from the vapor return line.

8.1.2.4 Concentration Measurement

Figure 17 illustrates equipment for concentration measurements of samples from manifolded vapor return lines.

8.1.3 Test Point 3 (Vent and/or Assist Processor)

Figure 18 emphasizes mass flux test locations for Test Point 3 (Vent and/or Assist Processor).

Test point 3 shall always be at the outlet from the vent riser. The operation of test equipment shall not interfere with the normal operation of any valve or vent.

8.1.3.1 Single Vent Volume Measurement

Figure 19 illustrates equipment for volume measurements of samples from a single vent at Test Point 3.

8.1.3.2 Manifolded Vents Volume Measurement

Figure 20 specifically illustrates metering equipment for volume measurements of samples from manifolded vents at Test Point 3.

8.1.3.3 Vent or Vents Concentration Measurement

Figure 21 illustrates equipment for concentration measurements of samples from a vent or vents.

8.1.4 Test Point 4 (Vapor Incinerator)

8.1.4.1 Incinerator Performance Specifications

Incinerator emissions shall be determined using the procedures of EPA M-2B, as outlined in this procedure, including any additional requirements provided below.

Any incinerator shall be evaluated and tested to establish:

- (1) a performance specification for carbon monoxide (CO) emissions and
- (2) performance specifications for other critical incinerator operating parameters per CP-201 § 3 which requires, in part:

The results of evaluation and testing of the system, documented in the certification test report, shall include:

(1) the identification of such critical system operating parameters.

- (2) the performance specifications for such critical system operating parameters, and
- (3) the specification of requirements for indicating gauges, detection devices, and alarms.

Challenge and failure mode testing shall be performed to establish system sensitivity to and performance specifications for the following variables:

- (1) storage tank ullage at start of liquid transfer
- (2) volume and volumetric rate of liquid transfer
- (3) number of nozzles in simultaneous use and
- (4) individual nozzle dispensing rates.

Compliance with the incinerator performance specifications shall be determined per CP-201, as applicable.

8.1.4.2 Incinerator Sampling Parameters

A preliminary evaluation of incinerator operation shall be conducted to determine data collection intervals for time and parameter magnitude for each parameter. Such intervals shall be chosen to provide calculated estimates of incinerator mass emissions factors which differ by no more than \pm 10% from actual, based on engineering judgment.

Data for each parameter shall be collected on such intervals.

Collect and record incinerator data for all of the parameters required to make a determination per EPA M-2B, with additional requirements for auxiliary fuel to expand the applicability of EPA M-2B:

V_{in}	=	total inlet volume entering vapor incinerator	
•••		(SCF)	

[HC]_{fuel} = hydrocarbon concentration of auxiliary fuel

(volume fraction)

[HC]_{out} = vapor incinerator outlet hydrocarbon

concentration (ppm)

[CO₂] = vapor incinerator outlet carbon dioxide

concentration (ppm)

[CO] = vapor incinerator outlet carbon monoxide

concentration (ppm)

Based on an engineering evaluation of a subject incinerator, the ARB Executive Officer may allow simplifying assumptions to be used in place of actual data collection. For example, for auxiliary fuel, it is often possible to use data from the fuel supplier.

8.1.4.3 Incinerator Visual Inspection

Visual Inspection. Any visible emissions except for steam, from vapor incinerators are an indication of poor combustion. An incinerator shall not emit air contaminants (not including moisture) in such a manner that the opacity of the emission is greater than 10 percent for a period or periods aggregating more than one minute in any 60 consecutive minutes; or greater than 40 percent opacity at any time. Should such visible emissions from the exhaust be detected, the control system is unacceptable and the problem must be corrected and an application made to the ARB Executive Officer for reconsideration for certification.

8.1.4.4 Incinerator Exhaust Sample Location

The vapor incinerator exhaust sample must be taken from the exhaust stack down-stream of the burner far enough to permit complete mixing of the combustion gases. For most sources, this point is at least eight stack dismeters downstream of any interference and two diameters upstream of the stack exit. There are many cases where these conditions cannot be met. The sample point shall be no less than one stack diameter from the stack exit and one stack diameter above the high point of the flame and be at a point of maximum velocity head. Vapor incinerator emissions shall be monitored for a 24 hour period beginning at the time of the first dispensing period.

8.1.4.5 Incinerator Inlet Sample Location

The vapor incinerator inlet sample and temperature and pressure measurements must be taken from a sample manifold attached to the inlet side of the volume meter which has been inserted at a break in the inlet line. The installation of test equipment shall not interfere with the normal operation of the vapor incinerator. Unaltered sample shall be returned to the sample manifold.

8.2 General Sampling Parameters

The test team shall collect and record frequent periodic or continuous measurements of the following sample gas variables shown in hexagon outlines in Figure 1:

HC = Hydrocarbon Concentration

CO = Carbon Monoxide Concentration

CO₂ = Carbon Dioxide Concentration

V = Volume

P = Pressure

T = Temperature

8.3 Other Sampling Parameters

Test Point 1 (Nozzle Sleeve)

Dispensed Fuel Vapor Pressure

Dispensed Fuel Volume

Test Point 2 (Vapor Return)

Dynamic Back-Pressure as Found

Test Point 3 (Vent or Vapor Processor)

Valve Cracking Pressure

Phase I Delivery Descriptions

Test Point 4 (Vapor Incinerator)

Design Operating Parameters

Actual Operating Parameters

9 QUALITY ASSURANCE / QUALITY CONTROL (QA/QC)

9.1 Analyzers

Perform a comprehensive calibration in the laboratory every six months. Check the analyzer with several known concentrations of calibration from reference cylinders to determine linearity.

9.2 Calibration Gases

Calibration gases are classified into three types:

(1) Standard Reference Materials

These are primary standards to which all other standards shall be traceable. For any substance for which no standard reference material is obtainable, a calibration gas of the highest level of accuracy and precision obtainable shall qualify as a standard reference material, subject to approval by the ARB Executive Officer.

A standard reference material, which normally is kept at a main laboratory, qualifies as an intermediate standard and as a working standard, too.

(2) Intermediate Standards

These are secondary standards which shall be assayed versus the corresponding NIST-SRM once every six months with a concentration difference which is no more than one percent of the results for the NIST-STAM. An intermediate standard container which does not meet its as requirement shall be taken out of service. To re-enter service, the intermediate standard container shall be recharged and meet its assay requirement.

An intermediate standard, which normally is kept at a branch laboratory or a shop, qualifies as a working standard, too.

(3) Working Standards

These are tertiary standards which shall be assayed versus the corresponding intermediate standard before every test with a concentration difference which is no more than one percent of the results for the intermediate standard. A working standard container which does not meet its assay requirement shall be taken out of service. To re-enter service, the working standard container shall be recharged and meet its assay requirement.

A working standard normally serves for field calibration and testing.

A "Certificate of Analysis" from the gas supplier can be submitted in the Certification Test Report as evidence of compliance with the specifications above; regardless of such certificate, the tester is ultimately responsible for satisfying the requirements given above in the event that a certificate is contradicted by subsequent analysis of the contents of a certified gas container.

All calibrations shall be performed with a calibration gas of at least working standard quality. Any cylinder is to be recharged or taken out of service when the cylinder pressure drops to 10 percent of the original pressure.

Information on calibration gas containers shall be entered into a permanent log identifying each container by serial number. Sufficient information shall be maintained to allow a determination of the compliance status of each calibration gas per these requirements; such information shall include for each container, but not be limited to each:

- (1) date put in service,
- (2) assay result, and
- (3) date taken out of service.

9.3 Volume Meters

Standard methods and equipment shall be used to calibrate the meters on an annual basis. The calibration curves are to be traceable to NIST standards.

10 RECORDING DATA

(1) Chain of Custody

Written data records must be kept during testing and kept by chain of custody.

(2) Necessary and Sufficient Data

Written data records must contain all information used to calculate and report final results.

(3) Reconciliation of Reported Results to Recorded Data

The final results must be verifiable by recalculation from the written data records.

(4) Permanent Records

These written data records must be kept permanently filed and available for use by the Executive Officer of the Air Resources Board when requested.

11 CALCULATING RESULTS

Note: In addition to other required calculations, vapor recovery system test results shall be calculated in units of pounds of hydrocarbon emitted per thousand gallons of fuel transferred for any results which are expressible in such units.

Calculate all efficiency results to the nearest 0.1%.

11.1 General Nomenclature

Figure 1 illustrates some parameters specified in the calculations.

11.1.1 Parameters

General parameters are listed below, other parameters are defined in the calculations or alternative procedures:

[HC] = hydrocarbon concentration (volume fraction),

V_m ≡ measured volume of gases and vapors,

P = pressure, and

T = temperature.

For any dispensing episode:

D = volume of liquid dispensed, and

 Δt = elapsed time of dispensing.

11.1.2 Subscripts

Subscripts shall be used to distinguish parameters and modes of measurement, e.g.:

P_(s,e,t) value of parameter "P" for subinterval "s" of dispensing episode "e" at test point "t".

Any or all of these subscripts may modify a parameter, and for consistency, subscripts will appear in the order given above, e.g.:

P_(e,t) = value of parameter "P" for dispensing episode "e" at test point "t"; and

P_t value of parameter "P" for an entire test at test point "t".

11.2 Standardization and Calibration of Parameters

11.2.1 Volume Standardization

Directly measured volumes (such as those directly measured for Test Points 1, 2, and 3) shall be standardized as follows:

$$V = V_{m} \left(\frac{528}{T} \right) \left[\frac{P_{b} + \left(\frac{P}{13.6} \right)}{29.92} \right]$$

where:

V = volume corrected to standard conditions (ft³).

 V_m = measured volume (ft³).

P_b = barometric pressure (in. Hg).

P = differential pressure in sample line (in. water gauge).

T = temperature of gas stream (°R).

11.2.2 Concentration

Each measured concentration of gas and vapor shall be corrected for any analyzer zero and/or span drifts and shall be expressed as a volume fraction (i.e. % or ppm).

11.2.3 Mass

Masses shall be calculated from calibration data and measurements as follows:

$$m = \left(\frac{MW}{385}\right) \times [HC] \times V$$

where:

m = mass (lb)

MW = molecular weight of calibration gas (lb/lb-mole)

385 = standard volume of one lb-mole at 528°R and 29.92 in. Hg

Note for manual data reduction: In general, $[HC]_{(e,t)}$ will stabilize to a steady value during a dispensing interval. If this is not the case, break $V_{(e,t)}$ into "s" subintervals and calculate:

$$m_{(e,t)} = \left(\frac{MW}{385}\right) \times \sum_{i=1}^{s} \left([HC]_{(s,e,t)} \times V_{(s,e,t)}\right)$$

11.3 Volume Calculations

11.3.1 Volume for Test Point 1 (Nozzle Sleeve)

This volume is directly measured and shall be standardized per § 11.2.1.

11.3.2 Volume for Test Point 2 (Vapor Return Line)

This volume is directly measured and shall be standardized per § 11.2.1.

11.3.3 Volume for Test Point 3 (Vent and/or Assist Processor)

This volume is directly measured and shall be standardized per § 11.2.1.

11.3.4 Volume for Test Point 4 (Incinerator)

Note the possibility for simplifying assumptions described in § 8.1.4.2.

11.3.4.1 Preliminary Incinerator Outlet Volume Calculations

Before calculating the vapor incinerator outlet volume, calculate the following preliminary values:

(1) inlet volume from the facility vapor space

Any inlet volume from the facility vapor space entering the vapor incinerator is directly measured and shall be standardized per § 11.2.1.

(2) inlet volume of auxiliary fuel

Any inlet volume from auxiliary fuel entering the vapor incinerator is directly measured and shall be standardized per § 11.2.1.

(3) total inlet volume entering vapor incinerator

$$V_{in} = V_{facility} + V_{fuel}$$

where:

(4) inlet hydrocarbon concentration

$$[HC]_{in} = \frac{(N [HC]_{facility} V_{facility}) + (N [HC]_{fuel} V_{fuel})}{V_{in}}$$

where:

11.3.4.2

Final Incinerator Outlet Volume Calculations

Calculate any vapor incinerator outlet volume using the following equation:

$$V_{out} = V_{in} \left[\frac{[HC]_{in}}{N [HC]_{out} + [CO_2] + [CO] - 300} \right]$$

where:

11.4 Calculations of Emissions During Idle Nozzle Episodes

To evaluate the possibility of a system eventually meeting the efficiency performance standard, these calculations must be completed before the Dispensing Facility Vent Calculations, the Individual Dispensing Episode Calculations, and the ultimate Efficiency Test Result Calculation.

Idle nozzle emissions of HC at a dispensing facility must be apportioned to each dispensing episode on a proportional basis of dispensed volume.

11.4.1 Total Idle Nozzle Emissions

Total idle nozzle emissions for all idle nozzle episodes:

m₁ = HC mass through mass flux area 1 (idle nozzle)

If the ARB Executive Officer determines that a portion of m1 is due to Phase I activity, then m1 may be diminished by that portion.

11.4.2 Apportioned Idle Nozzle Emissions

For any D.:

 $f(D_e,m_1)$ = the fraction of idle nozzle emissions assigned to each dispensing episode on a proportional basis of dispensed volume.

- (liquid volume dispensed);
 (all liquid volume dispensed during flux of m₁)
- $m_{(e,1)} = m_1 \times f(D_e, m_1)$

11.5 Dispensing Facility Vent Calculations

To evaluate the possibility of a system eventually meeting the efficiency performance standard, these calculations must be completed before the Individual Dispensing Episode Calculations and the ultimate Efficiency Test Result Calculation.

Vent emissions of HC at a dispensing facility must be apportioned to each dispensing episode on a proportional basis of dispensed volume.

11.5.1 Total Vent Emissions

Total vent emissions for all dispensing episodes:

 m_3 = HC mass through mass flux area 3 (vent)

If the ARB Executive Officer determines that a portion of m_3 is due to Phase I activity, then m_3 may be diminished by that portion.

11.5.2 Apportioned Vent Emissions

For any D_a:

- f(D_e,m₃) = the fraction of vent emissions assigned to each dispensing episode on a proportional basis of dispensed volume.
 - (liquid volume dispensed);
 (all liquid volume dispensed during flux of m₃)
- $m_{(e,3)} = m_3 \times f(D_{e}, m_3)$

11.6 Individual Dispensing Episode Calculations

Processor and incinerator emissions of HC at a dispensing facility must be apportioned to each dispensing episode on a proportional basis of dispensed volume. Use the same apportionment algorithm as for the vent emissions above.

The term "dispensing episode" is used here to generalize the applicability of these procedures.

Unless otherwise specified by the certification process, a dispensing episode starts with the removal of a nozzle from a dispenser and ends with the start of the next dispensing episode when the nozzle is removed again.

It is assumed that dispensing is into a vehicle fuel tank with a fillpipe test point and a vapor return line test point, but these calculations also apply to, for example, dispensing into surrogate tanks such as 55 gallon drums.

11.6.1 Mass through a Given Test Point

For any dispensing episode:

m_(e,t) = HC mass through a given test point

11.6.2 Individual Dispensing Episode Calculations

Each dispensing episode efficiency, Ee, is calculated from the mient:

$$E_{e} = \frac{m_{(e,2)} - [m_{(e,3)} + m_{(e,4)}]}{[m_{(e,2)} + m_{(e,1)}]} \times 100\%$$

where:

 $m_{(e,1)}$ = the mass flux through openings at the dispensing interface,

 $m_{(e,2)}$ = the mass flux through the vapor return line;

m_(e,3) = the mass flux through the vent and/or the assist processor; and

 $m_{(e,4)}$ = the mass flux through the vapor incinerator.

11.7 Efficiency Test Result Calculation

For the tested vapor recovery equipment, the efficiency test result, E, for this procedure is:

$$E = \sum_{1}^{n} \left[\frac{E_{\theta}}{n} \right]$$

where "n" is the number of dispensing episodes.

12 REPORTING RESULTS

Note: In addition to other required results, vapor recovery system test results shall be reported in units of pounds of hydrocarbon emitted per thousand gallons of fuel transferred for any results which are expressible in such units.

The following are required by § 10 RECORDING DATA:

- (1) Chain of Custody
- (2) Necessary and Sufficient Data
- (3) Reconciliation of Reported Results to Recorded Data
- (4) Permanent Records

Example report forms are provided in Forms 1 through 4 for generating written documents to meet these requirements. Other formats can be used; however, no test report shall be accepted or approved unless it contains at least the information specified in these forms.

All such forms must be written and submitted on acceptable media as specified by the the ARB Executive Officer on a case-by-case basis for each report.

In cases of conflict between hard copy and soft copy documents, the hard copy shall be presumed correct, unless a different determination is made by the ARB Executive Officer in special circumstances, which must be documented, in hard copy and soft copy, to the satisfaction of the ARB Executive Officer.

13 ALTERNATIVE TEST PROCEDURES

13.1 General Alternative Test Procedures

Test procedures, other than specified above, shall only be used if prior written approval is obtained from the ARB Executive Officer. In order to secure the ARB Executive Officer's approval of an alternative test procedure, the applicant is responsible for demonstrating to the ARB Executive Officer's satisfaction that the alternative test procedure is equivalent to this test procedure.

- (1) Such approval shall be granted on a case-by-case basis only. Because of the evolving nature of technology and procedures for vapor recovery systems, such approval shall not be granted in subsequent cases without a new request for approval and a new demonstration of equivalency.
- (2) Documentation of any such approvals, demonstrations, and approvals shall be maintained in the ARB Executive Officer's files and shall be made available upon request.

13.2 Test Procedures for Determining Incinerator Emissions

Incinerator emissions shall be determined using the procedures of EPA M-2B with the additional requirements provided in TP-205.2.

14 REFERENCES

This section is reserved for future specification.

15 EXAMPLE FIGURES AND FORMS

15.1 Figures

Each figure provides an illustration of an implementation which conforms to the requirements of this test procedure; other implementations which so conform are acceptable, too. Any specifications or dimensions provided in the figures are for example only, unless such specifications or dimensions are provided as requirements in the text of this or some other required test procedure.

Figure 1
Test Locations

Figure 2
Test Point 1 (Nozzle Sleeve)

Figure 3
Vehicle Leak Check Procedure (Nitrogen Pressurization)

Figure 4
Vehicle Leak Check Procedure: (Manual Compression)

Figure 5
Vehicle Leak Check Procedure: (Manual De-Compression)

Figure 6
Nozzle Sleeve Assembly: (Sectional View of Sleeve)

Figure 7
Nozzle Sleeve Assembly: (Axial View of Sleeve)

Figure 8
Nozzle Sleeve Assembly: (View of Sleeve on Nozzle)

Figure 9
Leak Check of Sleeve: (View of Combustible Gas Detector)

Figure 10
Leak Check of Sleeve: (View of Combustible Gas Detector in Use)

Figure 11
Nozzle Sleeve Measurements: (Volume Measurement)

Figure 12
Nozzle Sleeve Measurements: (Concentration Measurement)

Figure 13
Test Point 2 (Vapor Return Line)

Figure 14

Vapor Return Line (Volume Measurement, General)

Figure 15

Vapor Return Line (Volume Measurement, Single Vapor Return Line)

Figure 16

Vapor Return Line (Volume Measurement, Manifolded Vapor Return Lines)

Figure 17

Vapor Return Line (Concentration Measurement)

Figure 18

Test Point 3 (Vent and/or Assist Processor)

Figure 19

Single Vent (Volume Measurement)

Figure 20

Manifolded Vents (Volume Measurement)

Figure 21

Vent or Vents (Concentration Measurement)

15.2 Forms

Each form provides an illustration of an implementation which conforms to the requirements of this test procedure; other implementations which so conform are acceptable, too. Any specifications or dimensions provided in the forms are for example only, unless such specifications or dimensions are provided as requirements in the text of this or some other required test procedure.

Form 1

Chain of Custody

Form 2

Data

Form 3

Results

Form 4

Permanent Records

FIGURE 1 Test Locations

vant valvo (HC) hydrocarbon concentratio $\langle \mathtt{T} \rangle$ temperature (P) pressure

FIGURE 2 Test Point 1 (Nozzle Sleeve)

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FIGURE 3 Vehicle Leak Check Procedure (Nitrogen Pressurization)

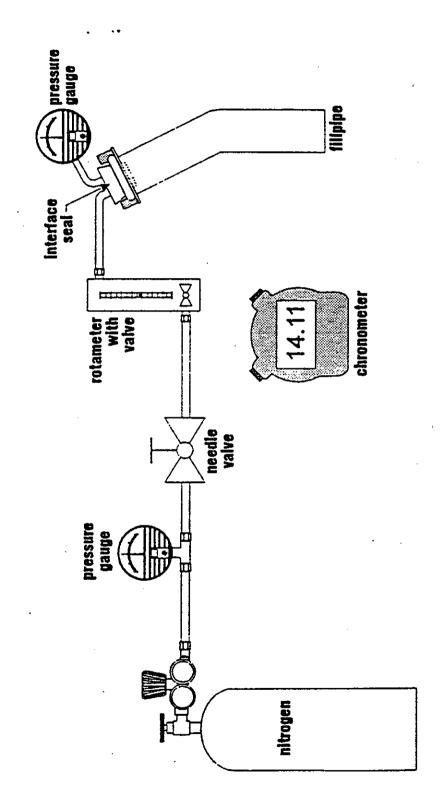
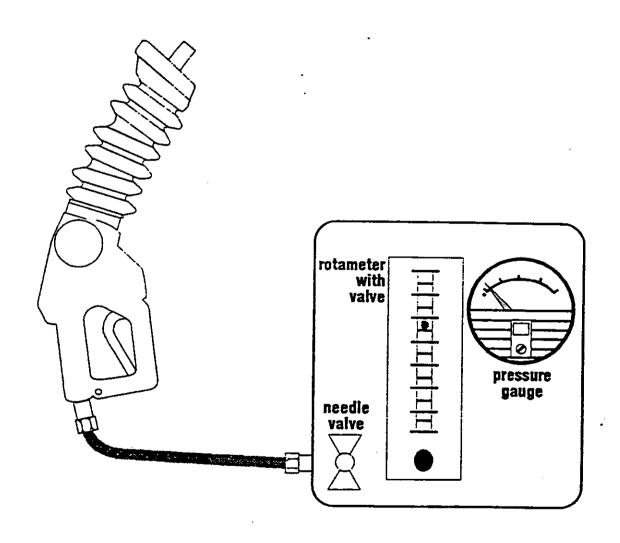
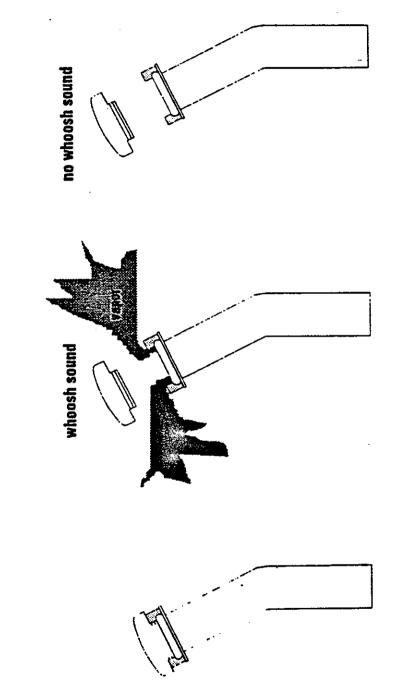


FIGURE 4 Vehicle Leak Check Procedure (Manual Compression)



TP 201.2 F.A/ B. CORDOVA TH

FIGURE 5 Vehicle Leak Check Procedure (Manual De-Compression)

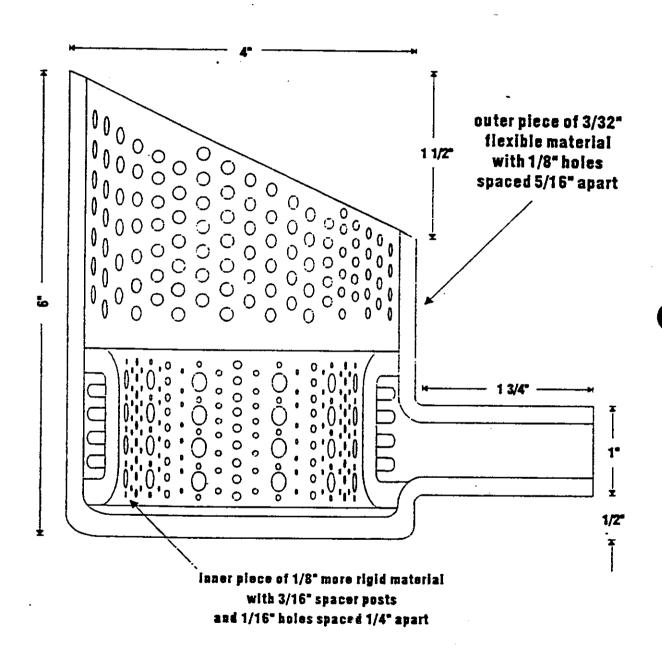


vent cap removed from fill-pipe

TP-201.2 page 43

vent cap on fill-pipe

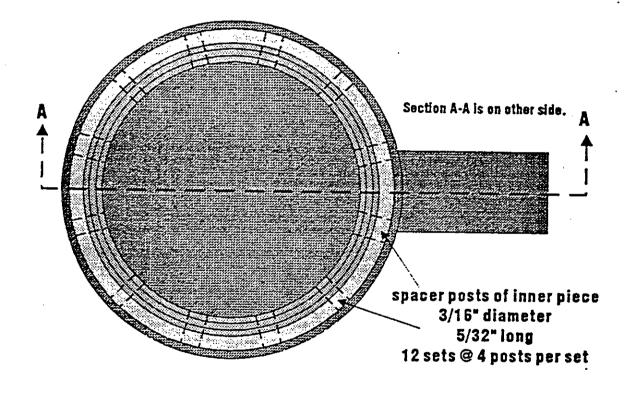
FIGURE 6 Nozzle Sleeve Assembly (Sectional View of Sleeve)



these dimensions are for example, and are not specifications

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FIGURE 7 Nozzle Sleeve Assembly (Axial View of Sleeve)



Materials must be resistant to breakdown by fuels and additives and easily bonded and repaired.

TP 201.2 F.7/ B. COROGYA 186

FIGURE 8 Nozzle Sleeve Assembly (View of Sleeve on Nozzle)

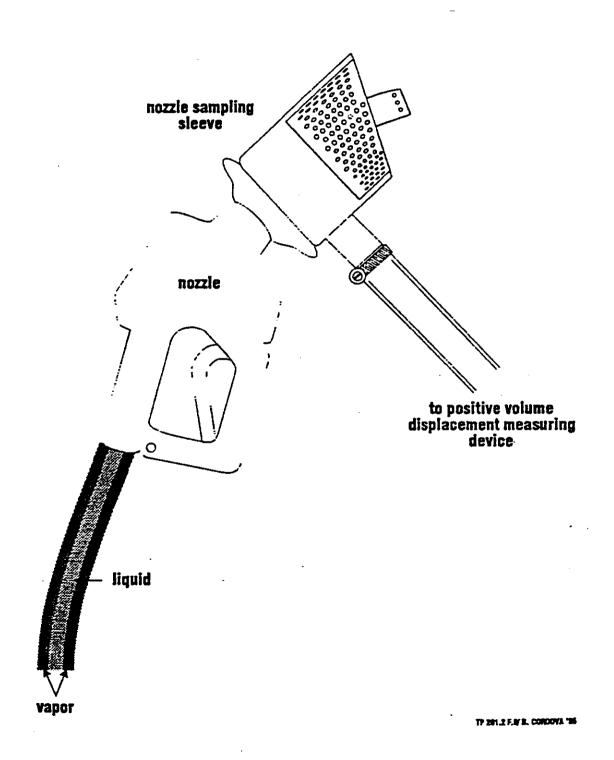


FIGURE 9 Leak Check of Sleeve (View of Combustible Gas Detector)

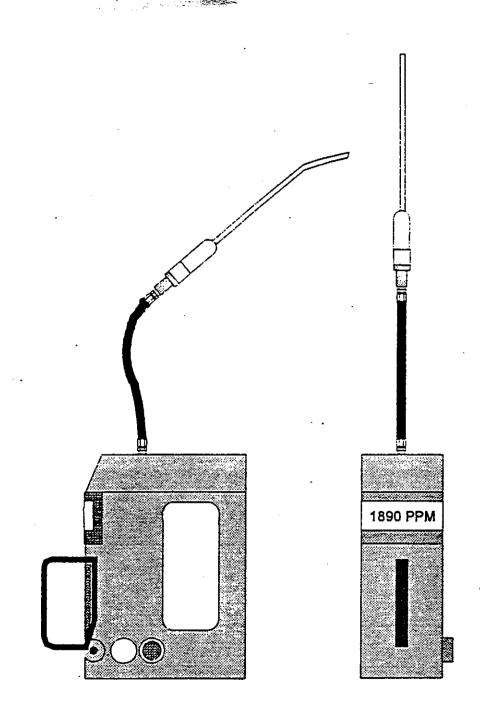
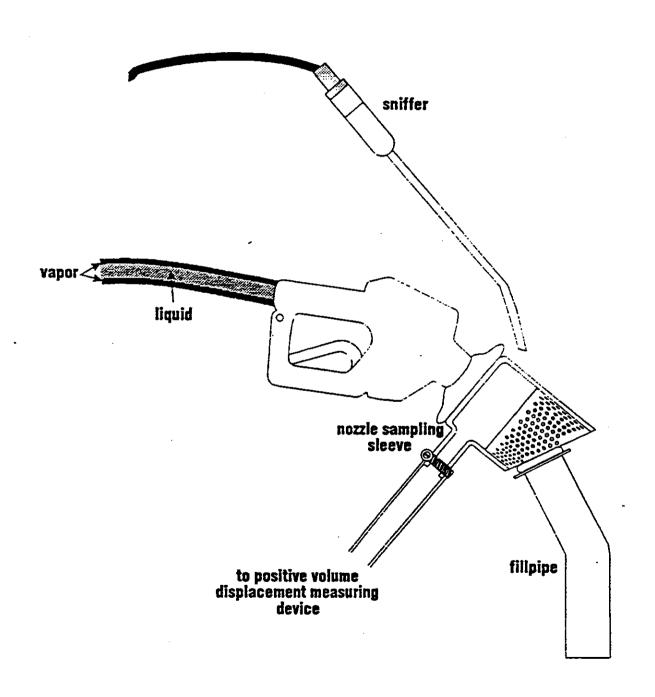


FIGURE 10 Leak Check of Sleeve (Combustible gas detector in Use)



TP 291,2 F.18/ R. COMDONA 195

FIGURE 11 Nozzle Sleeve Measurement (Volume Measurement)

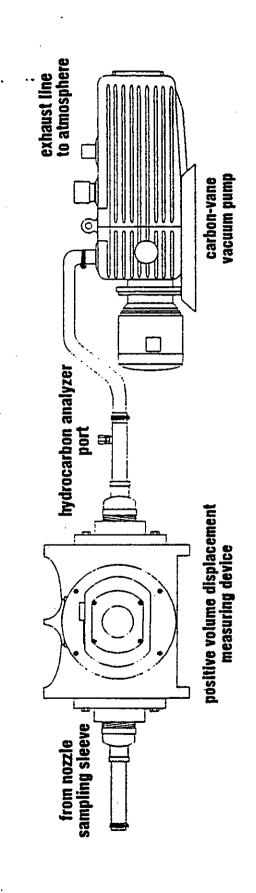
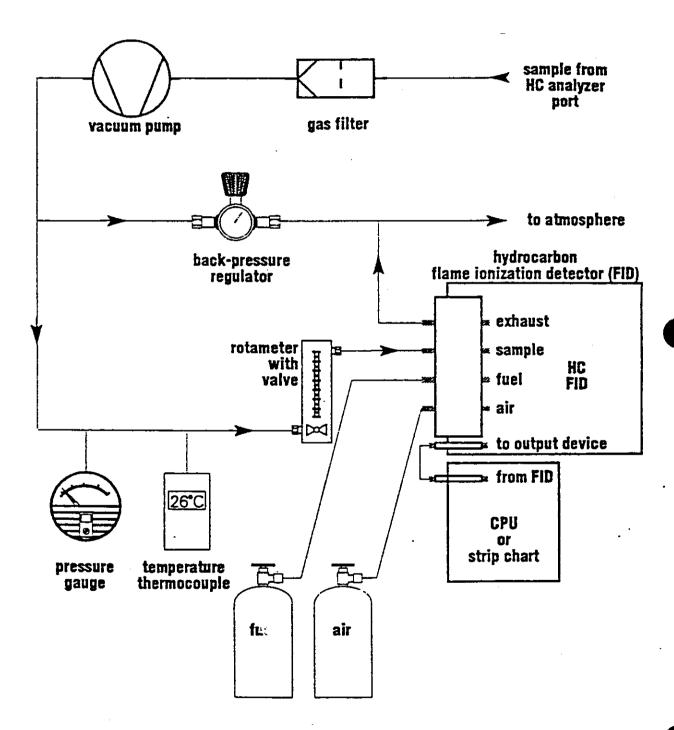


FIGURE 12
Nozzle Sleeve Measurements (Concentration Measurements)



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TP EST. F. T.V B. CORDOM, YE FIGURE 13 Test Point 2 (Vapor Return Line) HC hydrocarbon concentration
V volume
P pressure
T temperature underground tank

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FIGURE 14
Vapor Return Line (Volume Measurement, General)

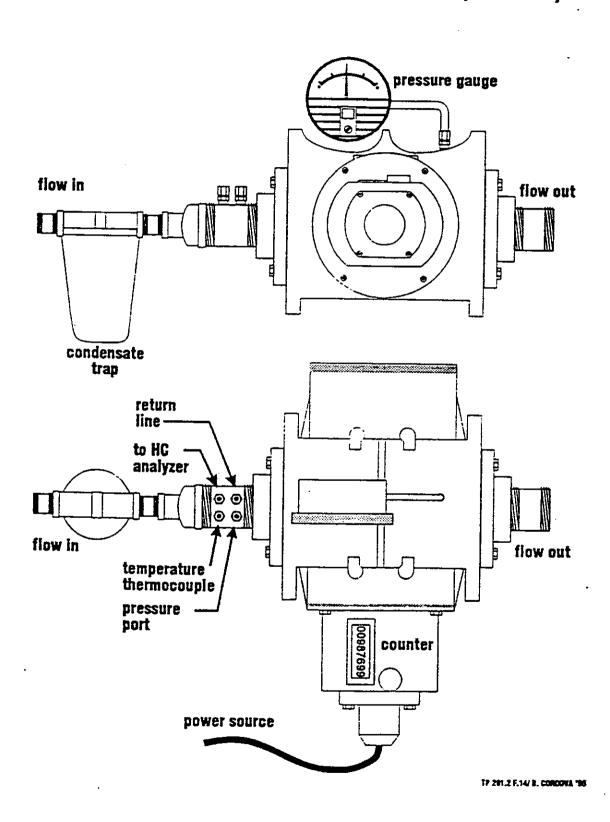


FIGURE 15 Vapor Return Line (Volume Measurement, Single Vapor Return Line)

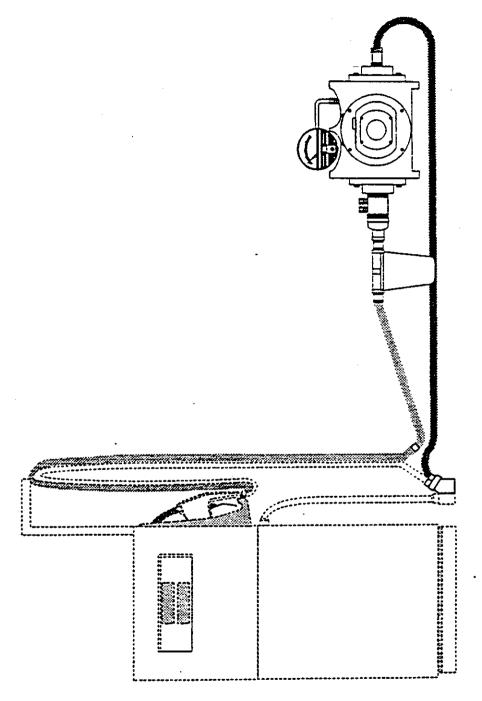


FIGURE 16
Vapor Return Line
(Volume Measurement, Manifolded Vapor Return Lines)

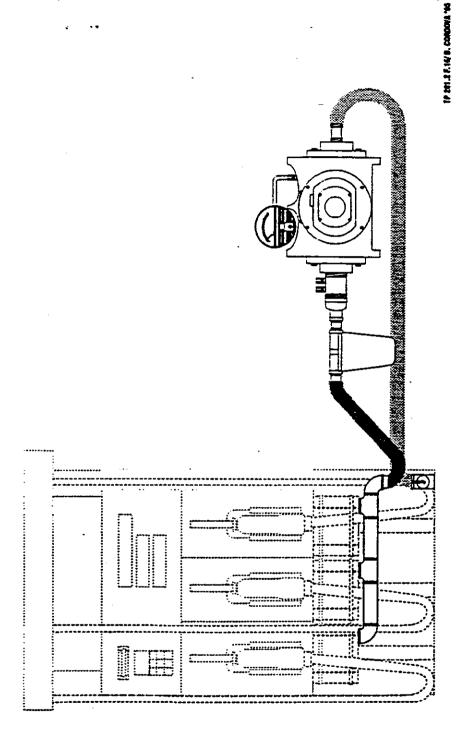
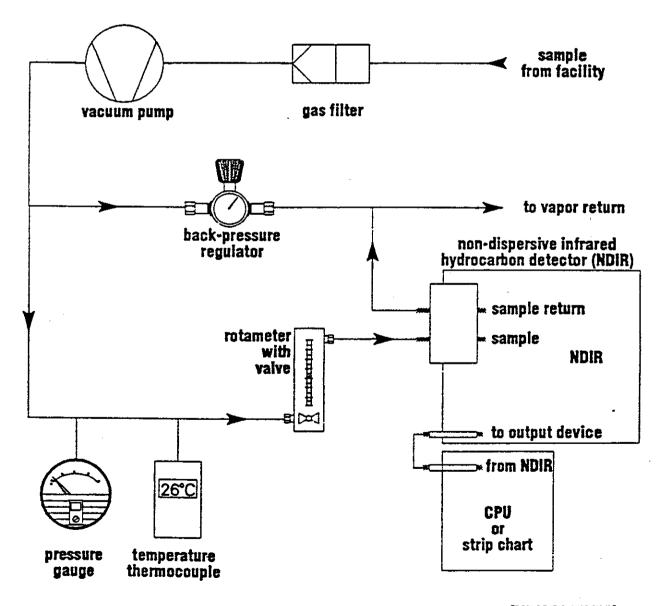


FIGURE 17
Vapor Return Line (Concentration Measurement)



TP 291.2 F.17/ B. CORDOVA '95

FIGURE 18 Test Point 3 (Vent and/ or Assist Processor)

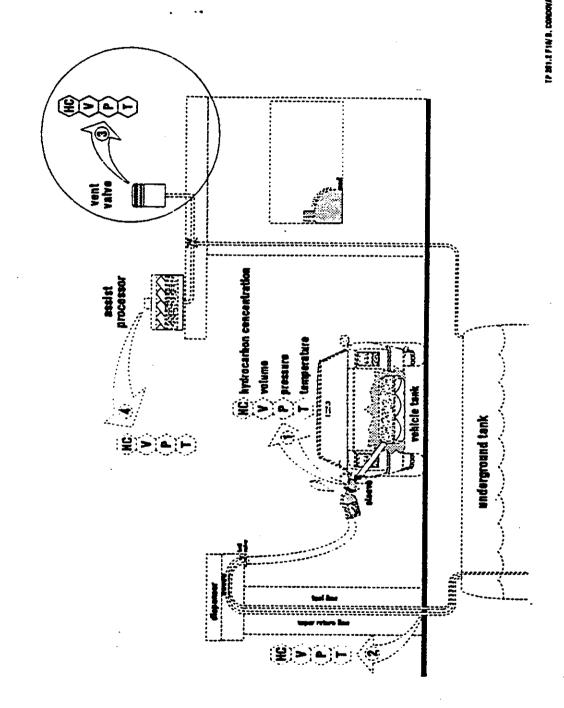


FIGURE 19 Single Vent (Volume Measurement)

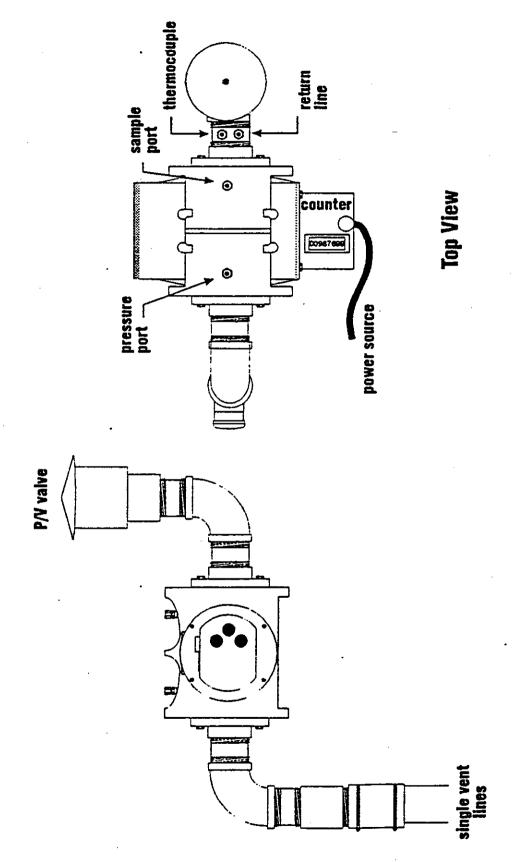
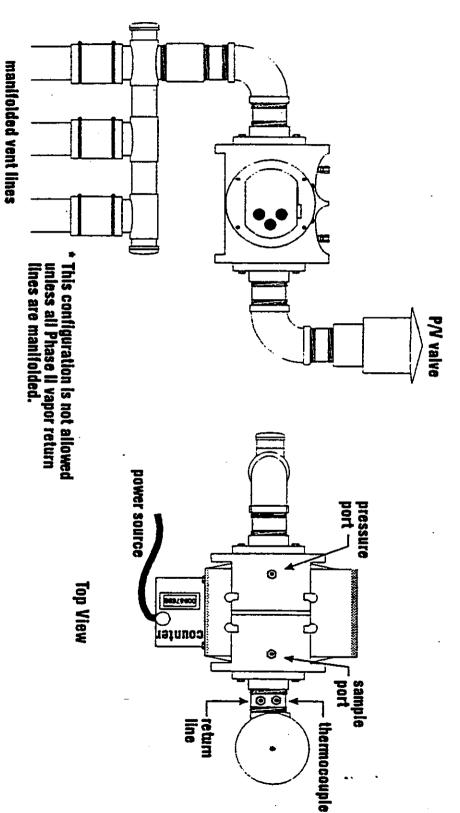
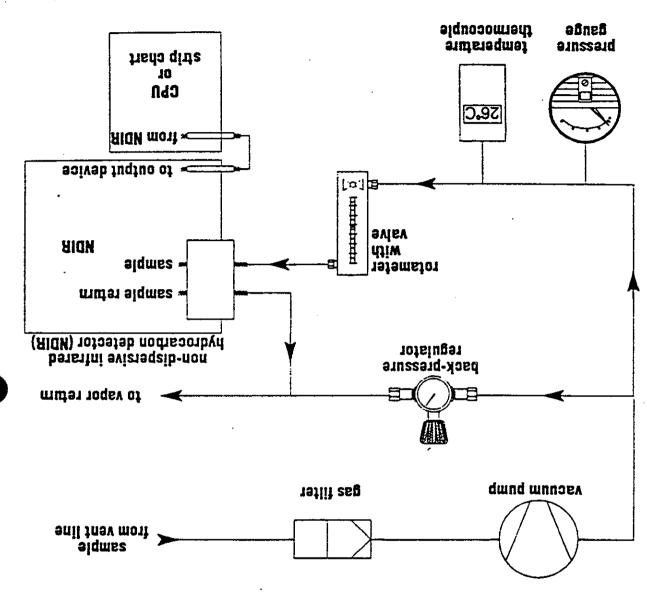


FIGURE 20 Manifolded Vents (Volume Measurement)

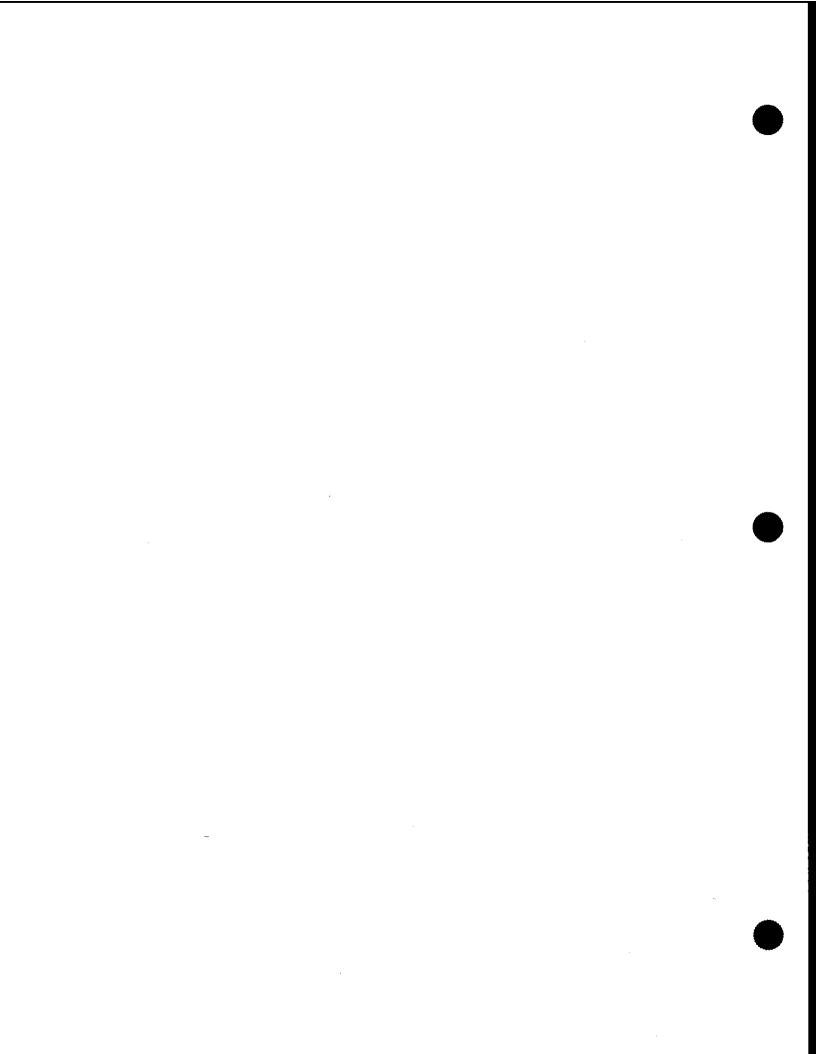


TP 301,2 F.34/ B, COMBOYA '98

Vent or Vents (Concentration Measurement)



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Form 1 Chain of Custody

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Phase II Notes	Q√ (ajmanyb)	(Joden)	p (vepor)	V _(Vapor)	[HC] ^(Astou)	# ebosid3	emiT
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			Location				System Description

Form S Data Test Point 3

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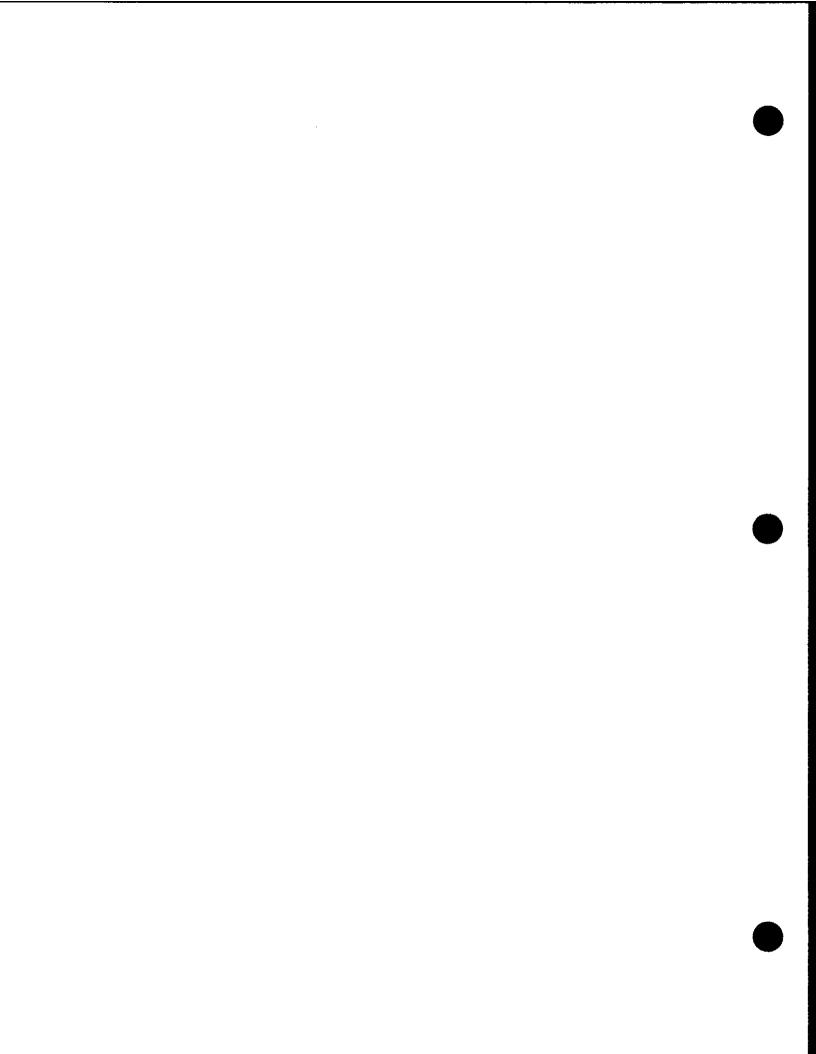
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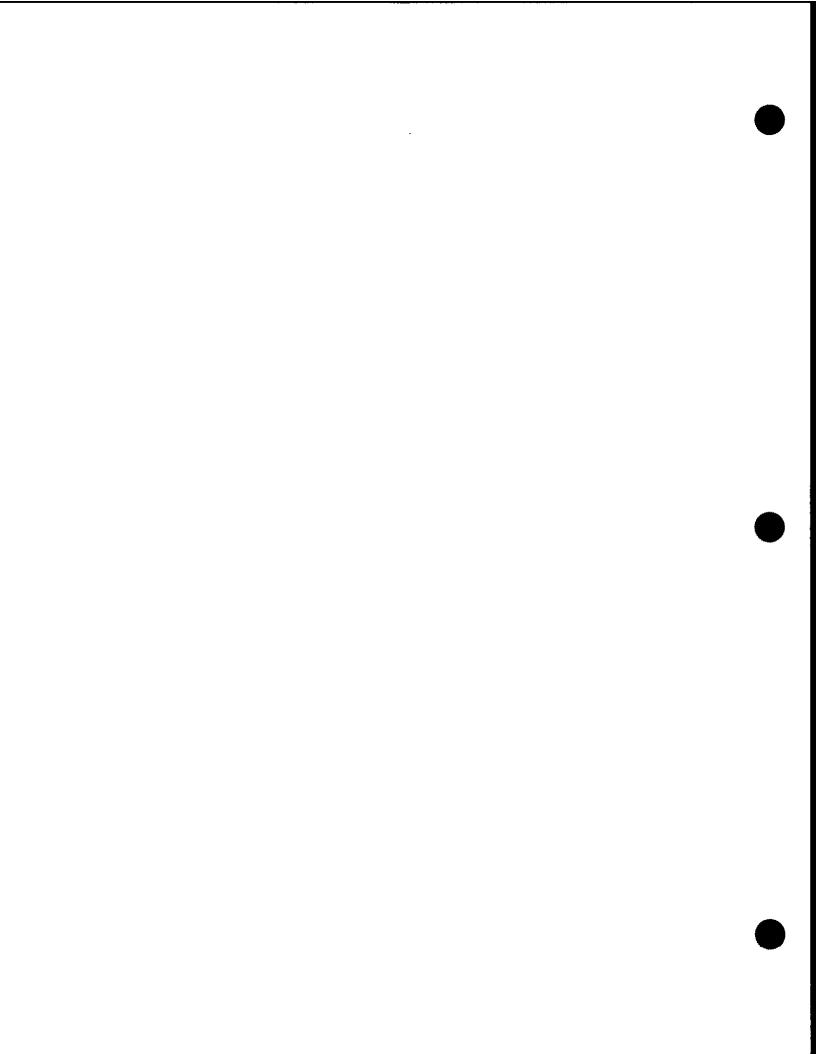
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Appendix B

TP-201.2A

DETERMINATION OF VEHICLE MATRIX FOR PHASE II VAPOR RECOVERY SYSTEMS OF DISPENSING FACILITIES



California Environmental Protection Agency



Air Resources Board

Vapor Recovery Test Procedure

TP-201.2A

Determination of Vehicle Matrix for Phase II Vapor Recovery Systems of Dispensing Facilities

Adopted: April 12, 1996

California Environmental Protection Agency Air Resources Board

Vapor Recovery Test Procedure

TP-201.2A

Determination of Vehicle Matrix for Phase II Vapor Recovery Systems of Dispensing Facilities

1 APPLICABILITY

Definitions common to all certification and test procedures are in:

D-200 Definitions for

Certification Procedures and

Test Procedures for

Vapor Recovery Systems

For the purpose of this procedure, the term "ARB" refers to the State of California Air Resources Board, and the term "ARB Executive Officer" refers to the Executive Officer of the ARB or his or her authorized representative or designate.

This test procedure can be used to determine the characteristics of a test fleet of vehicles which, when tested by other test procedures, can yield data representative of the total vehicle fleet.

2 PRINCIPLE AND SUMMARY OF TEST PROCEDURE

The sample of vehicles to be used in method TP-201.2 for testing vapor control systems shall be made up of vehicles representative of the on the road vehicle population in terms of vehicle miles traveled (VMT). This calculation procedure produces such a representative vehicle matrix. The distribution in terms of model year can be derived from the VMT portion of the calculated input to EMFAC. EMFAC is the ARB computer model for estimating on road motor vehicle emissions and is administered by the Technical Support Division of ARB. Distribution in terms of manufacturer can be derived from the number of registered vehicles for each make and model year which can be obtained from the Department of Motor Vehicles.

11 CALCULATING RESULTS

The vehicle makes and models and the number of vehicles per cell in the examples below are for illustration purposes only. More cells and other models and different numbers of vehicles per cell shall be included at the discretion of the ARB Executive Officer.

11.1 Vehicle Make

Obtain the number of registered vehicles by manufacturer and by model year from the Department of Motor Vehicles (DMV). The data shall resemble the following:

MODEL CHRY						
(e.g.)	SLER FORD	GM	TOYOTA	HONDA	OTHER	TOTAL
YEAR 1991 109,5 1990 138,4 etc.	•		218,577 203,156	191,174 189,973		1,577,886 1,668,708

11.4 VMT per Make for All Years

Sum the products of step 11.3 for each manufacturer. These sums represent the total VMT for each manufacturer. Select at least five manufacturers responsible for the highest VMT sums. These five (or more) manufacturers will be used to establish columns in the matrix. A last column called "Others" will include the vehicles from all other manufacturers.

11.5 Percentage of Vehicles for each Model Year

Determine the number of vehicles from each model year which are required by the 100-car matrix. To do this, convert the VMT fractions of step 11.2 to percents by multiplying by 100%. These percent numbers also equal the number of vehicles required in the 100-car test for each model year. For example if 10% of all VMT's are traveled by 1990 model vehicles, the 100-car matrix would include ten 1990 vehicles.

It is most accurate to maintain fractions through the calculations and round to whole vehicle numbers only at the last step of determining the matrix.

11.6 Percentage of Vehicles by Make for each Model Year

Obtain the fractions of registered vehicles by manufacturer for each model year. This shall be done for the five main manufacturers (step 11.4) and for the "Others" total. First, sum the numbers of registered vehicles of all manufacturers for each model year. Second, divide this sum into the registered vehicle numbers of each of the five main manufacturers and "Others" to get the desired fraction. For example, a recent calculation yielded:

Percentage of Registered Vehicles by Manufacturer for Each Model Year							
MODEL	CHRYSLER	FORD	GM	TOYOTA	HONDA	OTHERS	TOTAL
YEAR 1991	6.9	21.0	24.2	100	404	04.0	100.00/
1990 etc.	8.3	21.9 21.1	21.2 19.4	13.9 12.2	12.1 11.4	24.0 27.6	100.0% 100.0%

11.8 Vehicle Matrix

The vehicle matrix shall be constructed per the requirements of the certification procedure. Examples for two such requirements are given below:

11.8.1 Vehicle Cell Limits

The following example shows the results of constructing a vehicle matrix for August 1992 with a ten vehicle cell limit. Any other matrix with another cell limits, such as less than five vehicles per cell, shall be constructed in the same manner.

Combine the data into groups of model years to facilitate filling the matrix during the 100-car field test. Beginning with the current year, add previous years in succession until a maximum of ten vehicles accumulate in any cell. This group of model years will form the first row of cells. Repeat this process starting with the next preceding year to determine the group of years for the second row of cells. Repeat until all previous years combined yield less than 10 vehicles in any cell. This will normally require four rows of cells and the result will resemble the following table:

100 VEHICLE MATRIX AUGUST 1992							
MODEL YR	CHRYSLER	FORD	GM	TOYOTA	HONDA	OTHER	TOTALS
89-92	2	6	5	. 4	3	7	27
86-88	3	5	5	4	2	8	27
82-85	2	4	6	3	2	7	24
77-81	1	3	5	2	1	4	16
< 77	1	2	2	0	0	1	6
TOTALS	9	20	23	13	8	27	100

Be careful when rounding to whole numbers of vehicles. This can result in a matrix with slightly more or less than 100 vehicles. One can often determine the best place to add or subtract a vehicle by comparing the sums of rounded numbers and unrounded numbers for each row and column.

12 REPORTING RESULTS

This section is reserved for future specification.

13 ALTERNATIVE TEST PROCEDURES

Test procedures, other than specified above, shall only be used if prior written approval is obtained from the ARB Executive Officer. In order to secure the ARB Executive Officer's approval of an alternative test procedure, the applicant is responsible for demonstrating to the ARB Executive Officer's satisfaction that the alternative test procedure is equivalent to this test procedure.

- (1) Such approval shall be granted on a case-by-case basis only. Because of the evolving nature of technology and procedures for vapor recovery systems, such approval shall not be granted in subsequent cases without a new request for approval and a new demonstration of equivalency.
- (2) Documentation of any such approvals, demonstrations, and approvals shall be maintained in the ARB Executive Officer's files and shall be made available upon request.

14 REFERENCES

This section is reserved for future specification.

15 FIGURES

This section heading is not applicable to this procedure.

Appendix C

TP-201.2B

DETERMINATION OF FLOW VERSUS PRESSURE FOR EQUIPMENT IN PHASE II VAPOR RECOVERY SYSTEMS OF DISPENSING FACILITIES

State of California Air Resources Board

Vapor Recovery Test Procedure

PROPOSED TP-201.2B

Determination (Including Fugitive Emissions) of Efficiency of Phase II Vapor Recovery Systems of Dispensing Facilities

1 APPLICABILITY

A set of definitions common to all certification and test procedures is in:

D-200

Definitions for Certification Procedures and Test Procedures for Vapor Recovery Systems

For the purpose of this procedure, the term "ARB" refers to the State of California Air Resources Board, and the term "ARB Executive Officer" refers to the Executive Officer of the ARB or his or her authorized representative or designate.

A list of symbols and nomenclature used in this procedure is provided in the "REFERENCES" section.

1.1 General

This procedure applies to the determination of fugitive emissions from vapor recovery systems at dispensing facilities by direct measurement and modeling. This procedure applies to any fugitive vapor emissions associated with the dispensing of any fluid, although it is written to reflect application to the hydrocarbon (HC) vapors associated with the dispensing of gasoline.

1.2 Modifications

Modification of this procedure may be necessary for vapors and fluids other than the hydrocarbon vapors associated with the dispensing of gasoline.

Any modification of this method shall be subject to approval by the ARB Executive Officer.

2 PRINCIPLE AND SUMMARY OF TEST PROCEDURE

The purpose of this test procedure is to determine the fugitive emissions and the vapor recovery efficiency (including fugitive emissions) at a dispensing facility. Figures 1 through 3 are provided to illustrate some aspects of the principle and summary provided below. Figures are at the end of this document.

2.1 Principle

The mass flux of fugitive emissions from a dispensing facility is the product of the volumetric flow rate and the flow-weighted mass-per-volume concentration.

The volumetric flow rate is based upon data for pressure vs. time from the facility and data for flow vs. pressure from a model of the facility. The model flow vs. pressure data are provide a conversion for the facility pressure vs. time data to flow vs. time data.

The flow-weighted mass-per-volume concentration is based upon data for pressure vs. time from the facility, data for mass-per-volume concentration from specified components at the facility, and data for flow vs. pressure taken from specified components on a model of the facility. The model flow vs. pressure data provide a conversion for the facility pressure vs. time data to flow vs. time data for the specified components. The concentration and flow vs. time data for the specified components provide a basis for the flow-weighted mass-per-volume concentration.

2.2 Summary

As required to determine an emissions parameter and except where otherwise specified, the equipment and procedures specified in the following test methods shall be used.

EPA Method 18	Measurements of Gaseous Organic Compound Emissions by Gas Chromatography
EPA Method 25A	Determination of Total Gaseous Organic Compound Emissions Using a Flame Ionization Detector
EPA Method 25B	Determination of Total Gaseous Organic Compound Emissions Using a Nondispersive Infrared Analyzer

A detailed summary of this test procedure is provided in Figure 4.

3 BIASES AND INTERFERENCES

3.1 Static Pressure Performance, TP-201.3

Because the pressure data collection required by this test procedure are not valid unless the facility complies with the static pressure performance standard required by CP-201, the test procedure for static pressure performance, TP-201.3, shall be used to establish that the facility complies with the static pressure performance standard before and after the performance of this test procedure. If this requirement is not met, this test procedure must be performed again.

3.2 Interference by TP-201.2

Because the concurrent performance of TP-201.2 can interfere with the performance of this test procedure, this test procedure shall be performed during the initial 90 day reliability check required by CP-201.

4 SENSITIVITY, RANGE, AND PRECISION

This section is reserved for future specification. More developments are expected.

5 EQUIPMENT

5.1 Polished Stainless Steel Canisters

Polished stainless steel canisters are used to collect integrated samples for concentration analysis. Ancillary valves and flow meters are used to ensure that a low, constant sample flow rate is achieved which does not create sample artifacts by inducing flow in the vapor recovery system.

5.2 Hydrocarbon Analyzers

Integrated samples for concentration analysis must be analyzed both for total hydrocarbon concentration and for speciated hydrocarbon concentrations. Ancillary standard gases must approximate the range of species and concentrations expected for dispensing facility samples (100ppm_v).

6 CALIBRATION PROCEDURE

This section is reserved for future specification. More developments are expected.

7 PRE-TEST PROTOCOL

Perform TP-201.3, as required below.

8 TEST PROCEDURE

The purpose of this test procedure is to determine the fugitive emissions and the vapor recovery efficiency (including fugitive emissions) at a dispensing facility.

A list of symbols and nomenclature used in this procedure is provided in the "REFERENCES" section. Such symbols are used consistently in this section and in the "CALCULATING RESULTS" section. Figures 1 through 3 are provided to illustrate some aspects of the procedure required below. A detailed summary of this test procedure is provided in Figure 4. The test procedure follows:

Take and permanently record atmospheric pressure readings which are chronologically coordinated with all other readings required by this procedure:

where:

(P)_{201.2A, atm} = atmospheric pressure, "WC.

8.1 Find $(Q_{201.3, \Sigma F})$ at $(P_{201.3}) = 5\text{"WC}_g$ for the total facility

Perform TP-201.3.

Find (Q_{201.3}, $_{\Sigma F}$) at (P_{201.3}) = 5"WC_g for the total facility at the test conditions of TP-201.3

Warning: Do not proceed with this test procedure until TP-201.3 has been performed and $(Q_{201.3,\;\Sigma F})$ has been calculated as specified the "CALCULATING RESULTS" section.

where:

 $(Q)_{201.3, \Sigma F} \equiv \text{flow, ft}^3/\text{min}$

 $(P)_{201.3}$ = initial pressure, "WC_a

 $(V)_{201.3}$ = facility vapor volume, gal

 $(\Delta t)_{201.3}$ = elapsed time, min

 $(\Delta P)_{201.3}$ = absolute value of pressure change, "WC

(P)_{201.3, atm} = atmospheric pressure, "WC.

where:

$$(P)_{2F}$$
 = pressure, "WC_a

Assemble and secure the required equipment.

Perform calibrations as required.

Take and permanently record pressure readings at one minute intervals for two weeks.

Perform QA/QC as required.

8.3 Find (T) and [HC_{m/v}] vs. (t) for specified facility components

Find (T) and $[HC_{m/v}]$ for representative episodes for vapors behind the facility components which have vapor valves:

Take samples at the following sample locations and determine values of the variables listed for each sampling location:

1F (Idle Nozzle Valves)

$$[HC_{m/v}]_{1F}$$
 = hydrocarbon concentration, lb_m/ft^3

$$[HC_{v/v}]_{1F}$$
 = hydrocarbon concentration, volume fraction

$$[HC_{MW}]_{1F}$$
 = hydrocarbon molecular weight, lb_m/lb -mole

$$(T)_{1F}$$
 = temperature, °F

2F (Overfill Drain Valves)

$$[HC_{m/v}]_{2F}$$
 = hydrocarbon concentration, Ib_m/ft^3

$$[HC_{v/v}]_{2F}$$
 = hydrocarbon concentration, volume fraction

$$[HC_{MW}]_{2F}$$
 = hydrocarbon molecular weight, Ib_m/Ib -mole

$$(P)_{2F} \equiv \text{pressure, "WC}_n$$

 $(T)_{2F}$ = temperature, °F

3F (Vent Valves)

 $[HC_{m/v}]_{3F}$ = hydrocarbon concentration, Ib_m/ft^3

 $[HC_{v/v}]_{3F}$ = hydrocarbon concentration, volume fraction

[HC_{MW}]_{3F} ≡ hydrocarbon molecular weight, lb_m/lb-mole

 $(T)_{3F}$ = temperature, ${}^{\circ}F$

 $(z)_{3F-2F}$ = typical height above 2F, in

 $(P)_{2F-3F}$ = typical pressure below $(P)_{2F}$, "WC

8.3.1 Test representative episodes

Unless the tester documents episodes of facility operations as more representative to the satisfaction of the ARB Executive Officer, the following episodes chosen by the ARB Executive Officer shall be tested as representative of facility operations:

- two hours of facility operations including the one hour of maximum dispensing to vehicle tanks,
- (2) two hours of facility operations including the one hour of minimum dispensing to vehicle tanks, and
- (3) two hours of facility operations following the delivery of liquid to a facility storage tank from a cargo tank.

Find (T) and [HC_{m/v}] vs. (t) for each episode for each sample location

For each component sample location: 1F (Idle Nozzle Valve), 2F (Overfill Drain Valve), and 3F (Vent Valve):

- (1) Assemble and secure the required equipment.
- (2) Perform calibrations as required.
- (3) Take samples for two hours per EPA Method 18.
- (4) Perform QA/QC as required.

For each episode and sample, perform a speciated gas chromatographic analysis for hydrocarbons per EPA Method 18 and find values for $[HC_{m/v}]_{(s)} \ lb_m/ft^3$ (hydrocarbon concentration, pounds per cubic foot, at sample location "s").

Warning: Do not proceed with this test procedure until the $[HC_{m/v}]_{(s)}$ have been calculated as specified the "CALCULATING RESULTS" section.

1.0 =
$$\sum_{i=1}^{n} ([HC_i])_{\{t,e,s\}}$$

where:

"n" = the number of hydrocarbon species (or species categories).

8.3.3 Calculate P_(2F-3F)

Warning: Do not proceed with this test procedure until the $P_{\{2F-3F\}}$ have been calculated as specified the "CALCULATING RESULTS" section.

8.4 Calculate $(P_{\Sigma F})$ vs. (t) for the facility

Note:

 $(P_{\Sigma F})_{(t)} \equiv \text{zero for any system with no vent valve}$

Warning: Do not proceed with this test procedure until the $(P_{\Sigma F})$ have been calculated as specified the "CALCULATING RESULTS" section.

8.5 Model values of specified variables for the facility and components

Design and validate a model of the facility and find values for specified variables of the facility, using the model.

8.5.1 Design the model

Design the model so that it can:

8.5.1.1 Reproduce (Q) vs. (P) for the total facility

Establish (Q) vs. (P) similitude with the total facility by reproducing (Q) vs. (P) for the model at the facility test conditions of TP-201.3.

8.5.1.2 Include components of the facility

Include components of the facility which have vapor valves.

Secure randomly sampled units of such components and avoid testing components which are not expected to be representative of components which ultimately will be installed at new installations of the system tested.

8.5.2 Find (Q)_{M, ΣF} vs. (P)_{M, ΣF} < 5"WC_a

Find (Q)_{M, ΣF} vs. (P)_{M, ΣF} for the total facility at other (P)_{M, ΣF} < 5"WC_g, using the model.

Volumetrically standardize results at 528°R and 1 atm:

where:

$$(P)_{M, \Sigma F} \equiv \text{pressure, "WC}_{\alpha}$$

$$(Q)_{M, \Sigma F} \equiv flow, ft^3/min$$

$$(T)_{M, \Sigma F}$$
 = temperature, °F

Warning: Do not proceed with this test procedure until the $(Q)_{M, \Sigma F}$ vs. $(P)_{M, \Sigma F}$ have been standardized as specified the "CALCULATING RESULTS" section.

8.5.3 Find $(Q)_{M, \Sigma F}$ vs. (t) for the total facility

Find (Q)_{M, ΣF} vs. (t) for the total facility, using the results of §8.4 and §8.5.2.

Note:

 $(Q)_{M, \Sigma F} \equiv \text{zero for any system with no vent valve}$

8.5.4 Find the flow-weighted average, $[HC_{avg m/v}]_{\Sigma F}$

Find the flow-weighted average, $[HC_{avg\ m/v}]_{\Sigma F}$, for the total facility, using the model.

8.5.4.1 Find (Q)_M vs. (P)_{M,} < 5"WC_g for the components

Find (Q) vs. (P) for the components at the facility which have vapor valves at other (P)_{M, ΣF} < 5"WC_g, using the model of the facility.

8.5.4.2 Find (Q)_(M) vs. (t)

for the components

Find $(Q)_{(M)}$ vs. (t) for the components at the facility which have vapor valves, using the results of §8.4 and §8.5.4.1.



(Q)_{M, (s)} = zero for any system with no vent valve

8.5.4.3 Calculate the flow-weighted average, $[HC_{avg m/v}]_{\Sigma F}$

Calculate the flow-weighted average, $[HC_{avg\ m/v}]_{\Sigma F}$, for the total facility by finding (for each of the components at the facility which have vapor valves) the summation of $(Q) \times [HC_{m/v}] \times (t)$ divided by the summation of $(Q) \times (t)$.

Warning: Do not proceed with this test procedure until [HC $_{\rm avg~m/v}]_{\Sigma F}$ has been calculated as specified the "CALCULATING RESULTS" section.

$$\frac{[HC_{\text{avg m/v}}]_{\Sigma F} \#}{ft^{3}} = \frac{\sum_{s=1}^{n} \sum_{i=1}^{n} \left[\frac{[HC_{\text{m/v}}]_{(i, s)} \#}{ft^{3}} \times \frac{(Q)_{(i, s)} ft^{3}}{\min} \times (t)_{(i, s)} \min \right]}{\sum_{s=1}^{n} \sum_{i=1}^{n} \left[\frac{(Q)_{(i, s)} ft^{3}}{\min} \times (t)_{(i, s)} \min \right]}$$

where:

"s" \equiv the valve component type corresponding to sample location 1F through 3F

"i" = the "ith" valve of valve component type "s" at the facility

"n" \equiv the number of valves of valve component type "s" at the facility.

8.6 Find (HC)_{ΣF} emitted from the facility

See "CALCULATING RESULTS" section.

8.7 Find (HC)_{ΣF} emissions factors for the facility

See "CALCULATING RESULTS" section.

8.8 Find the vapor recovery efficiency (including fugitive emissions) for the facility

See "CALCULATING RESULTS" section.

9 QUALITY ASSURANCE / QUALITY CONTROL (QA/QC)

Two collocated canister samples shall be taken at each sample location to allow for the calculation of relative error.

10 RECORDING DATA

This section is reserved for future specification. More developments are expected.

11 CALCULATING RESULTS

Note: In addition to other required calculations, vapor recovery system test results shall be calculated in units of pounds of hydrocarbon emitted per thousand gallons of fuel transferred for any results which are expressible in such units.

A list of symbols and nomenclature used in this procedure is provided in the "REFERENCES" section. Such symbols are used consistently in this section and in the "TEST PROCEDURE" section. Figures 1 through 3 are provided to illustrate some aspects of the procedure required below.

A detailed summary of this test procedure is provided in Figure 4.

11.1 Volume Standardization

Volumes shall be standardized as follows:

$$(V_{std}) \text{ ft}^3 = (V_m) \times \frac{(P_m) \text{ "WC}}{407 \text{ "WC}} \times \frac{528 \text{ }^{\circ}\text{R}}{(T_m) \text{ }^{\circ}\text{R}}$$

where:

$$(V_{std})$$
 = volume corrected to standard conditions, ft³

$$(V_m)$$
 = measured volume, ft³

$$(P_m)$$
 = measured pressure, "WC

$$(T_m)$$
 = measured temperature, ${}^{\circ}R$.

11.2 Find $(Q_{201.3, \Sigma F})$ at $(P_{201.3}) = 5\text{"WC}_g$ for the total facility

Find ($Q_{201.3, \Sigma F}$) at ($P_{201.3}$) = 5"WC_g for the total facility at the test conditions of TP-201.3:

$$\frac{(Q)_{201.3, \Sigma F} \text{ ft}^3}{\text{min}} = \frac{(V)_{201.3}}{(\Delta t)_{201.3}} \times \frac{(\Delta P)_{201.3}}{(P)_{201.3, atm}} \times \frac{1 \text{ ft}3}{7.481 \text{ gal}}$$

11.3 Find [HC_{m/v}] vs. (t) for each episode for each sample location

$$\frac{[HC_{m/v}]_{(s)} \#}{ft^3} = \sum_{i=1}^{n} \left[[HC_{v/v}]_{(i)} \times \frac{(MW)_{(i)} \#}{\#mole} \times \frac{\#mole}{385 \text{ ft}^3} \times \frac{407 \text{ "WC}}{(P)_{(201.2A, \text{ atm})} \text{ "WC}} \times \frac{(T)_{(i)} \text{ °R}}{528 \text{ °R}} \right]_{(s)}$$

where:

1.0 =
$$\sum_{i=1}^{n} ([HC_i])_{(t,e,s)}$$

"n" = the number of hydrocarbon species (or species categories).

11.4 Calculate P(2F-3F)

$$(P)_{2F-3F}$$
 "WC = $(z)_{3F}$ in $\times \frac{([HC_{m/v}]_{3F} - AIR DENSITY_{m/v}) \#}{ft^3} \times \frac{ft^3}{62.4 \#}$

where:

AIR DENSITY_{m/v} is calculated by application of the Ideal Gas Equation.

11.5 Calculate $(P_{\Sigma F})$ vs. (t) for the facility

$$(P_{\Sigma F})$$
 "WC = $(P)_{2F}$ "WC - $(P)_{2F-3F}$ "WC

11.6 Calculate the flow-weighted average, $[HC_{avg m/v}]_{\Sigma F}$

Calculate the flow-weighted average, $[HC_{avg\ m/v}]_{\Sigma F}$, for the total facility by finding (for each of the components at the facility which have vapor valves) the summation of (Q) \times $[HC_{m/v}] \times$ (t) divided by the summation of (Q) \times (t).

$$\frac{[HC_{avg\ m/v}]_{\Sigma F} \#}{ft^3} = \frac{\sum_{s=1}^{n_s} \sum_{i=1}^{n_i} \left[\frac{[HC_{m/v}]_{(i,\ s)} \#}{ft^3} \times \frac{(Q)_{(i,\ s)} ft^3}{min} \times (t)_{(i,\ s)} min \right]}{\sum_{s=1}^{n_s} \sum_{i=1}^{n_i} \left[\frac{(Q)_{(i,\ s)} ft^3}{min} \times (t)_{(i,\ s)} min \right]}$$

where:

"s" = the valve component type corresponding to sample location 1F through 3F

"n_s" = the number of valve component types at the facility

"i" \equiv the "ith" valve of valve component type "s" at the facility

"n;" = the number of valves of valve component type "s" at the facility.

11.7 Find (HC)_{ΣF} emitted from the facility

Find $(HC)_{\Sigma F}$ emitted from the facility during any specified time interval by finding $(Q)_{\Sigma F} \times [HC_{m/v}]_{avg} \times (t)$ for the total facility.

11.8 Find $(HC)_{\Sigma F}$ emissions factors for the facility

Find $(HC)_{\Sigma F}$ emissions factors for the facility for specified combinations of test parameters, e.g.: $(HC)_{\Sigma F}/(t)$, $(HC)_{\Sigma F}/(t)$, etc.

11.9 Find the vapor recovery efficiency (including fugitive emissions) for the facility

Find (HC)_{ΣF} for the duration of TP-201.2

$$m_5 = \sum_{i=1}^{n} (Q_{\Sigma F})_{(i)} \times ([HC_{m/v}]_{avg})_{(i)} \times (t)_{(i)}$$

where:

m5 = fugitive mass emitted for all time intervals of TP201.2, lb_m

 $t_{(i)}$ = "ith" time interval of TP201.2, min

Fugitive emissions shall be considered as being released from sample location (5) for the purposes of the efficiency calculation below, using the results of TP-201.2:

11.9.1.1 Apportioned Fugitive Emissions

Fugitive emissions of HC at a dispensing facility must be apportioned to each dispensing episode ($D_{\rm e}$) on a basis proportional to dispensed volume.

For any Da:

$$\Phi(D_e)$$
 = $\frac{\text{(liquid volume dispensed)}_e}{\text{(all liquid volume dispensed during flux of ms)}}$

where:

 $\Phi(D_e)$ = the mass fraction of fugitive emissions assigned to each dispensing episode on a proportional basis of dispensed volume.

"e" = "eth" dispensing episode of TP-201.2

 $m_{(e, 5)} = m_5 \times \Phi(D_e)$

where:

m_(e, 5) = the mass of fugitive emissions assigned to each dispensing episode on a proportional basis of dispensed volume

11.9.1.2 Individual Dispensing Episode Calculations

Unless otherwise specified by the certification process, a dispensing episode starts with the removal of a nozzle from a dispenser and ends with the start of the next dispensing episode when the nozzle is removed again.

It is assumed that dispensing is into a vehicle fuel tank with a fillpipe test point and a vapor return test point, but these calculations also apply to, for example, dispensing into surrogate tanks such as 55 gallon drums.

11.9.1.3 Mass through a Given Test Point

For any dispensing episode:

 $m_{(e,s)}$ = HC mass through a given sampling location (or apportioned fugitive mass emissions)

11.9.2 Individual Dispensing Episode Calculations

Each dispensing episode efficiency, E_e , is calculated from the $m_{(e,s)}$:

$$E_{e} = \frac{m_{(e,2)} - [m_{(e,3)} + m_{(e,4)} + m_{(e,5)}]}{[m_{(e,2)} + m_{(e,1)}]} \times 100\%$$

11.10 Find the vapor recovery efficiency (including fugitive emissions) for the facility

For the tested vapor recovery equipment, the efficiency test result, E, for this procedure is:

$$E = \sum_{e=1}^{n} \left(\frac{E_e}{n} \right)$$

where "n" is the number of dispensing episodes.

12 REPORTING RESULTS

Note: In addition to other required results, vapor recovery system test results shall be reported in units of pounds of hydrocarbon emitted per thousand gallons of fuel transferred for any results which are expressible in such units.

This section is reserved for future specification. More developments are expected.

13 ALTERNATIVE TEST PROCEDURES

Test procedures, other than specified above, shall only be used if prior written approval is obtained from the ARB Executive Officer. In order to secure the ARB Executive Officer's approval of an alternative test procedure, the applicant is responsible for demonstrating to the ARB Executive Officer's satisfaction that the alternative test procedure is equivalent to this test procedure.

- (1) Such approval shall be granted on a case-by-case basis only. Because of the evolving nature of technology and procedures for vapor recovery systems, such approval shall not be granted in subsequent cases without a new request for approval and a new demonstration of equivalency.
- (2) Documentation of any such approvals, demonstrations, and approvals shall be maintained in the ARB Executive Officer's files and shall be made available upon request.

14 REFERENCES

14.1 Symbols and Nomenclature

14.1.1 Format

Abbreviations

Abbreviations are defined to provide shorter descriptions for variables, subscripts, and units. Abbreviations used throughout this procedure for fugitive emissions paths are:

paths through "closed" idle nozzle check valves

2F = paths through "closed" overfill drain valves

3F = paths through "closed" vent valves

4F = other paths

ΣF

all paths

Variables

Variables are defined with an abbreviation in parentheses or square brackets:

()

general variable

[]

concentration variable

Subscripts

Subscripts are defined to distinguish test variables and test modes, e.g.:

(P)_(t, e, s)

value of parameter "(P)" for time interval "(t)" of testing episode "(e)" for sample location "(s)".

Any or all of these subscripts may modify a parameter and, for consistency, subscripts are defined in the order given above, e.g.:

(P)_(e, s)

value of parameter "(P)" for testing episode "(e)" for sample location "(s)".

 $(P)_{(s)}$

value of parameter "(P)" for entire test for sample location "(s)".

Subscripts are defined to distinguish types of concentrations, e.g.:

[(C)]_(v/v)

concentration variable (volume fraction)

 $[(C)]_{(m/v)}$

concentration variable (mass/volume)

[(C)]_(MW)

concentration variable (mass/mole)

Units

Units are defined and used without punctuation. The definition of each variable ends with a comma, followed by abbreviated units for such variable. Abbreviations for units used in this procedure are:

°R

■ degrees Rankine

ft

= feet

gal ≡ gallons

in = inches

"WC = inches of water column (pressure)

"WC_g = inches of water column (gauge pressure)

min = minutes

 385ft^3 = volume of lb-mole (at P_{std} and T_{std})

 $lb_m f$ = pounds (force)

 $lb_m = pounds (mass)$

Ib-mole ≡ pound mole

407"WC \equiv standard pressure (P_{std})

 $528^{\circ}R$ = standard temperature (T_{std})

Values

Values are substituted for variables and are used without punctuation.

14.1.2 General Test Variable

(P)_{201.2A, atm} = atmospheric pressure, "WC

14.1.3 Facility Test Variables

Figures 1 and 2 illustrate some facility sample locations and fugitive emissions test variables.

Static Pressure Performance Results (TP-201.3)

 $(\Delta P)_{201.3}$ = absolute value of pressure change, "WC

(P)_{201.3, atm} = atmospheric pressure, "WC

 $(\Delta t)_{201.3}$ = elapsed time, min

 $(V)_{201.3}$ = facility vapor volume, gal

 $(Q)_{201.3, \Sigma F} \equiv \text{flow, ft}^3/\text{min}$

 $(P)_{201.3}$ = initial pressure, "WC_a

1F (idie Nozzle Valves)

 $[HC_{m/v}]_{1F}$ = hydrocarbon concentration, lb_m/ft^3

[HC_{viv}]_{1F} = hydrocarbon concentration, volume fraction

 $[HC_{MW}]_{1F}$ = hydrocarbon molecular weight, lb_m/lb -mole

(T)_{1F} = temperature, °F

2F (Overfill Drain Valves)

 $[HC_{m/v}]_{2F}$ = hydrocarbon concentration, lb_m/ft^3

 $[HC_{v/v}]_{2F}$ = hydrocarbon concentration, volume fraction

 $[HC_{MW}]_{2F}$ = hydrocarbon molecular weight, Ib_m/Ib -mole

 $(P)_{2F}$ = pressure, "WC_a

 $(T)_{2F}$ = temperature, °F

3F (Vent Valves)

 $[HC_{m/v}]_{3F}$ = hydrocarbon concentration, Ib_m/ft^3

 $[HC_{v/v}]_{3F}$ = hydrocarbon concentration, volume fraction

 $[HC_{MW}]_{3F}$ = hydrocarbon molecular weight, lb_m/lb -mole

 $(T)_{3F}$ = temperature, ${}^{\circ}F$

 $(z)_{3F-2F}$ = typical height above 2F, in

 $(P)_{2F-3F}$ = typical pressure below $(P)_{2F}$, "WC

14.1.4 Model Test Variables

Figure 3 illustrates some model sample locations and test variables.

ΣF (Total Facility)

[HC]_{M, ΣF, avg m/ν}

hydrocarbon concentration (flow-weighted average mass/volume), lb_m/ft³

 $(P)_{M, \Sigma F}$ = pressure, "WC_a

 $(Q)_{M, \Sigma F} = flow, ft^3/min$

 $(T)_{M, \Sigma F}$ = temperature, °F

1F (Idle Nozzie Valves)

(P)_{M, 1F}

pressure, "WC_a

(Q)_{M. 1F}

flow, ft³/min

(T)_{M, 1F}

= temperature, °F

2F (Overfill Drain Valves)

(P)_{M. 2F}

≖ pressure, "WC_a

(Q)_{M, 2F}

≡ flow, ft³/min

(T)_{M, 2F}

3F (Vent Valves)

(P)_{M, 3F}

≖ pressure, "WC_α

(Q)_{M, 3F}

 \equiv flow, ft³/min

(T)_{M. 3F}

= temperature, °F

14.1.5 Test Procedure Results

This section is reserved for future specification. More developments are expected.

14.2 TP-201.2

14.3 TP-201.3

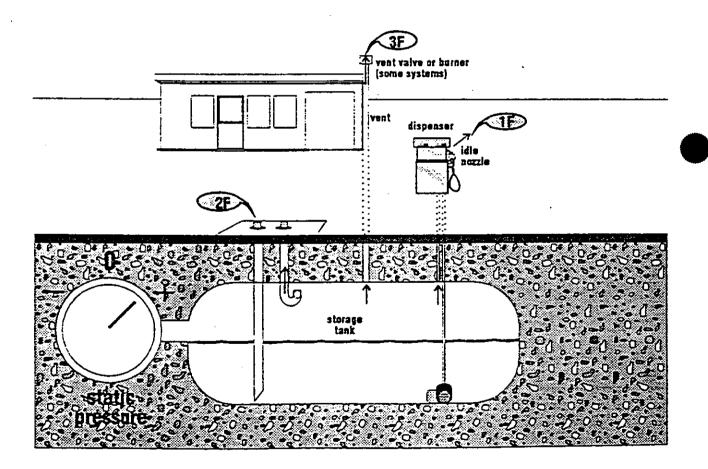
15 FIGURES

Figures are attached.

Figure 1

Examples of Locations of Equipment to be Tested

1F "closed" idle nozzle check valves 2F "closed" overfill drain valves 3F "closed" vent valves



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FIGURE 2 Examples of Equipment to be Tested

idle nozzles overfill drains vents

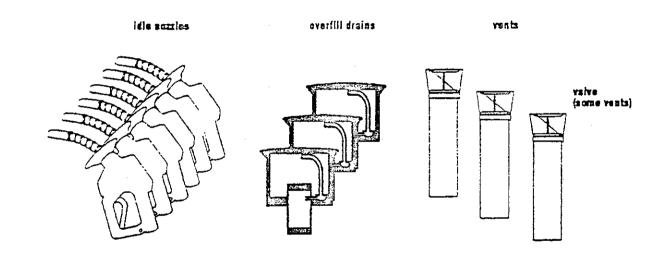


FIGURE 3 Example of a Bench Test

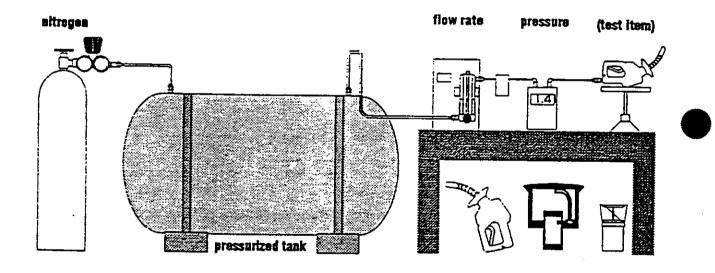


FIGURE 4 (continued)

Summary of Test Procedure

- (4) Design and validate a model of the facility and find values for specified variables of the facility, using the model.
 - (a) Design the model so that it can:
 - (i) establish (Q) vs. (P) similitude with the total facility by reproducing (Q) vs. (P) for the model at the facility test conditions of TP-201.3 and
 - (ii) include components of the facility which have vapor valves.
 - (b) Find (Q) vs. (P) for the total facility at other (P), using a model of the facility.
 - (c) Find (Q) vs. (t) for the total facility, using the results of (2) and (4)(b).
 - (d) Find the flow-weighted average, $[HC]_{avg\ m/v}$, for the total facility.
 - (i) Find (Q) vs. (P) for the components at the facility which have vapor valves at other (P), using the model of the facility.
 - (ii) Find (Q) vs. (t) for the components at the facility which have vapor valves, using the results of (2) and (4)(d)(i).
 - (iii) Find the flow-weighted average, $[HC]_{avg\ m/v}$, for the total facility by finding (for each of the components at the facility which have vapor valves) the summation of $(Q) \times [HC_{m/v}] \times (t)$ divided by the summation of $(Q) \times (t)$, using the results of (3) and (4)(d)(ii).
- (5) Find (HC) emitted from the facility during any specified time interval by finding (Q) \times [HC_{m/v}]_{avg} \times (t) for the total facility.
- (6) Find (HC) emissions for the facility for specified combinations of test parameters, e.g.:
 - (a) (HC)/(t),
 - (b) (HC)/(V), etc.
- (7) Find the vapor recovery efficiency (including fugitive emissions) for the facility.

FIGURE 4

Summary of Test Procedure

The tester must determine values for the following variables of a dispensing facility using conventional principles of sampling and analysis:

(HC) = hydrocarbon mass

 $[HC]_{m/v}$ = hydrocarbon concentration (mass/volume)

[HC]_{avg m/v} = hydrocarbon concentration (flow-weighted average mass/volume)

(P) ≡ pressure

(Q) = volumetric flow rate

(T) = temperature

(t) = time

(V) ≡ volume.

Then, the tester must find correlated values of a model of the dispensing facility using conventional principles of modeling, sampling, and analysis.

Finally, the tester must find (HC) emissions factors for the facility for specified combinations of test parameters.

In more detail, the principle of this procedure is to:

- (1) Find (Q) vs. (P) for the total facility at the test conditions specified in TP-201.3.
- (2) Find (P) vs. (t) for the total facility for representative episodes.
- (3) Find (T) and $[HC]_{m/v}$ for representative episodes for vapors behind the facility components which have vapor valves.

(continued on next page)

Appendix D 1998 MODELS EQUIPPED WITH ORVR SYSTEMS

Onboard Refueling Vapor Recovery 1998 MODEL YEAR CERTIFICATES Light-Duty Vehicles October 3, 1997 with

	EPID	RPID	BPID						RACER	TRACER WAGON
	CHRYSLER: CONCORDE; DODGE; INTREPID	CHRYSLER: CONCORDE; DODGE: INTREPID	CHRYSLER: CONCORDE; DODGE: INTREPID	DAEWOO: LANOS	DAEWOO: LANOS	DAEWOO: NUBIRA	FORD: ESCORT WAGON	Added: ford: escort	LINCOLN-MERCURY: TRACER WAGON, TRACER FORD: ESCORT, ESCORT WAGON	ADDED: LINCOLN-MERCURY: TRACER, TRACER WAGON
09-24-97	09-15-97 TLEV	09-15-97 TLEV	09-24-97	08-27-97 TLEV	08-22-97 Tlev	09-23-97 TLEV	07-30-97	07-30-97	07-30-97	07-31-97
ਜ	.	н	Ħ	~ 4	Ħ	п	H		н	
REVISED	WCRXV0195V20/WCRXR0101G1G 20LDV21	WCRXV0165V20/WCRXR0101G1G 20LDV22	WCRXVO2.7VB0/WCRXR0101GBG 20LDVZ3	WDWXV01.6D01/WDWXR0095A0L 178LDV01	WDWXVO1.5501/WDWXR0095A0L 178LDVO5	WDWXV02,0D01/WDWXR0095AOL 1761DV06	WEMXVOZ.OBEA/WEMXROOBOBAE	REVISED	WFMXVO2.OATA/WFMXROOBOBOE 3OLDV21	REVISED
		CARB A-9-382		DAENGO	•.		WORD		CARB A-10-736	

SUBARU: FORESTER AWD

09-02-97

-

WFJXV02.5CAC/WFJXR01251BB

660LDV08

WFJXV02.SDCD/WFJXR01251BB

CARB A-2-105-1-A

660LDV09

SUBARU: FORESTER AWD

09-02-97

TLRV

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EPA
1998 MODEL YEAR CERTIFICATES
Light-Duty Vehicles
with
Onboard Refueling Vapor Recovery

October 3, 1997

FORD: CROWN VICTORIA, CROWN VICTORIA POLICE FORD: CROWN VICTORIA, CROWN VICTORIA POLICE LINCOLN-MERCURY: GRAND MARQUIS, TOWN CAR LINCOLN-MERCURY: GRAND MARQUIS, TOWN CAR LINCOLN-MERCURY: TRACER, TRACER WAGON SUBARU: LEGACY AND, LEGACY WAGON AWD, SUBARU: LEGACY AWD, LEGACY WAGON AWD, SUBARU: LEGACY AWD, LEGACY WAGON AWD, SUBARU: LEGACY AND, LEGACY WAGON AWD, IMPREZA AWD, IMPREZA WAGON AWD IMPREZA AWD, IMPREZA WAGON AWD IMPREZA AWD, IMPREZA WAGON AWD IMPREZA AMD, IMPREZA WAGON AWD FORD: EXCORT, ESCORT WAGON; 07-31-97 09-25-97 10-02-97 05-08-97 05-08-97 05-08-97 05-08-97 TLEV TLEV -H H -1 4 WPMXV02.OALA/WFMXR0080BOB WFMXVO4.6ABA/WFMXR0115BAE WPMXV04.6AAA/WFMXR0115BAE WEJXV02.2AAA/WEJXR01251BB WFJXV02.2AAA/WFJXR01251BB WFJXV02.2BCB/WFJXR01251BB WFJXV02.2BCB/WFJXR01251BB FUJI-LDV-98-05 FUJI-LDV-98-04 FUJI-LDV-98-07 FUJI-LDV-98-6 30LDV25 30LDV27 30LDV29 CARB A-10-737 CARB A-2-104 FUJI

EPA
1998 MODEL YEAR CERTIFICATES
Light-Duty Vehicles
with
Onboard Refueling Vapor Recovery
October 3, 1997

GENERAL MOTORS

				-
	WGMXV01.9001/WGMXR0080902 GM-LDV-98-11	- 1	05-27-97	SATURN: SC, SL, SW
	WGMKV01.9003/WGMKR0080902 GM-LDV-98-12	г	05-27-97	SATURN: SC, SL, SW
CARB A-6-784	WGMXV01.9002/WGMXR0080902 GM-LDV-98-13	н	05-27-97 TLKV	SATURN: SC, SL, SW
CARB A-6-785	WGMXV01.9004/WGMXR00B0902 GM-LDV-98-14	н	06-05-97 Tlev	SATURN: SC, SL, SW
	WGMXVO3.1041/WGMXR0124912 GM-LDV-98-17	н	26-91-90	CHEVROLET: MALIBU; PONTIAC: GRAND PRIX; BUICK: CENTURY; OLDSMOBILE: CUTLASS
	WGMXVO3.8050/WGMXR0133910 GM-LDV-98-19	=	06-13-97	BUICK: RIVIERA, PARK AVENUB
	WGMXV03.8050/WGMXR0133918 GM-LDV-98-20	Ħ	06-13-97	PONTIAC: GRAND PRIX; BUICK: REGAL
	WGMXV04,6065/WGMXR0133910 GM-LDV-98-23	Ħ	06-19-97	CADILLAC: SEVILLE, OLDSMOBILE: AURORA
CARB A-6-797-A	WGMXV04.6066/WGMXR0133910 GM-LDV-26	н	07-01-97	CADILLAC: SEVILLE; OLDSMOBILE: AURORA
CARB A-6-796-A	WGMXV03.8051/WGMXR0133918	гſ	07-10-97	PONTIAC:GRAND PRIX, BUICK:REGAL

1998 MODEL YEAR CERTIFICATES
Light-Duty Vehicles
with
Onboard Refueling Vapor Recovery
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	BUICK: RIVIBRA, PARK AVENUE	CHEVROLET: MALIBU; OLDSMOBILE: CUTLASS; PONTIAC: GRAND PRIX; BUICK CENTURY	BUICK: PARK AVENUE	PONTIAC: GRAND PRIX; OLDSMOBILE: INTRIGUE; BUICK: REGAL	BUICK: PARK AVENUE	PONTIAC: GRAND PRIX; OLDSMOBILE: INTRIGUE; BUICK: REGAL	CHEVROLET: MALIBU; OLDSMOBILE: CUTLASS	CHEVROLET: MALIBU; OLDSMOBILE: CUTLASS	CHEVROLET; MALIBU; OLDSWOBILE; CUTLASS
	07-10-97	07-10-97	07-17-97	07-17-97	07-17-97 TLEV	07-17-98 TLEV	07-17-97	07-17-97 Teby	09-09-97 Tlev
	Ħ	н	н	=	н	ਜ	н	ri	ਜ
GM-LDV-98-28	WGMXVO3.8051/WGMXR0133910 GM-LDV-98-29	WGMXV03.1043/WGMXR0124912 GM-LDV-98-31	WGMXV03.8047/WGMXR0133910 GM-LDV-98-35	WGMXVO3.8047/WGMXR0133918 GM-LDV-98-36	WGMXV03.8048/WGMXR0133910 GM-LDV-98-37	WGMXVO3.8048/WGMXRO133918 GM-LDV-98-38	WGMXVO2.4024/WGMXR0124912 GM-LDV-98-39	WGMKV02,4026/WGMKR0124912 GN-IDV-98-40	WGMXVO3.1044/WGMXRO124912 40LDV42
	CARB A-6-796-A	CARB A-6-792-A			CARB A-6-794	CARB A-6-794		CARB A-6-789	CARB A-6-828-A

1998 MODEL YEAR CERTIFICATES Light-Duty Vehicles with

Onboard Refueling Vapor Recovery October 3, 1997

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1998 MODEL YRAR CERTIFICATES
Light-Duty Vehicles
with
Onboard Refueling Vapor Recovery
October 3, 1997

HYUNDAI: ELANTRA

HYUNDIA: TIBURON

HYUNDAI: ELANTRA

HYUNDAI: TIBURON

KIA: SEPHIA

	1 08-21-97 TLEV	1 08-21-97 TLEV	1 08-27-97	1 08-27-97	1 09-11-97
	WHYXVO1.82GM/WHYXRO13421E 265LDV07	WHYXV02.02GM/WHYXR013421E 265LDV08	WHYXVO1.81EL/WHYXRO1341EE 265LDV09	WHYXVO2.01TB/WHYXR01341EE 265LDV10	Wemkvol.8aol/Wemkrolooaol 338ldvol
HYUNDAI	CARB A-254-54	CARB A-254-55			KIA MOTORS .

08-11-97 MAZDA: 626	08-15-97 MAZDA: 626	08-11-97 MAZDA: 626
-80	08-	90
н	н	н
WTKXVO2.OVBA/WTKXRO125BFA 560ldvo6	WTKXVO2.OVDM/WTKXRO125BFA 560LDV07	WTKKVOZ.5VB2/WTKKRO125BFA 560LDVO8
MAZDA MOTOR	CARB A-16-224	

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1998 MODEL YEAR CERTIFICATES
Light-Duty Vehicles
with
Onboard Refueling Vapor Recovery
October 3, 1997

MERCEDES BENZ	WMBXVO3.2GNB/WMBXRO155MYN 200LDV09	н	09-25-97 LEV	MERCEDES-BENZ: E320
	WMBXVO2.3GSU/WMBXRO115MYY 200LDV10	н	09-25-97	Mercedes-Benz: SLK230
	WMBXVO5.OGNB/WMBXRO155MYY 200LDV11	H	09-29-97 TLEV	MERCEDES-BENZ: SL500
	WMBXVO6.0GNU/WMBXRO155MYY 200LDV12	н	09-29-97	MERCEDES-BENZ: SL600
MITSUBISHI (196 & 490) WDS.	£ 490) WDSXV02,4G2G/WDSXR016511A 196LDV04	ਜ	06-24-97 LKV	DIAMOND STAR MOTORS/MITSUBISHI: ECLIPSE CONVERT.
•	WDSXVO2.4GPG/WDSXRO16511A 196LDV06	H	06-24-97	DIAMOND STAR MOTORSIMITSUBISHI BCLIPSE CONVERT.
••••••	WDSXV02.5GFG/WDSXR016511A 196LDV08		07-02-97	DIAMOND STAR MOTORS/CHRYSLER: SEBRING, DODGE AVENGER
CARB A-292-40	WDSXV02.5G1G/WDSXR016511A 196LDV09 REVISED DATE 07-03-97	н	07-02-97 TLEV 08-12-97	CHRYSLRR: SEBRING; DODGE; DODGE AVENGER

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1998 MODEL YEAR CERTIFICATES
Light-Duty Vehicles
with
Onboard Refueling Vapor Recovery
October 3, 1997

Валв	WSAXVO2.3ND1/WSAXVOOBOYD1 470LDV02	Ħ	09-11-97	SAAB: SAAB 900, SAAB 900 CONVERTIBLE
	MSAXVO2.OTD1/WSAXVOOBOXD1 470LDV03	Ħ	. 09-11-97	SAAB: SAAB 900, SAAB 900 CONVERTIBLE
SUZUKI	WSKXV1.30CNA/WSKXROO85RMA SUZUK-LDV-98-04	н	06-12-97	SUZUKI: SWIFT; CHEVROLET:METRO
CARB-A-259-61	WSKXV1.30LNA/WSKXROO85RMA SUZUK-LDV-98-06	H	07-03-97 LEV	SUZUKI: SWIFT; CHEVROLET: METRO
TOYOTA	WTYXVO2.2XBA/WTYXRO135AK1 570LDV01	ч	07-15-97	TOYOTA; CAMRY
CARB A-14-319	WTYXVO2.2GXB/WTYXR0135AK1 570LDV02	H	07-31-97	TOYOTA: CAMRY
	WTYXVO1.8XBA/WTYXRO115AK1 570LDV05	H	07-25-97	TOYOTA: COROLLA
CARB A-14-318	WTYXVO1.8DXB/WTYXR0115AK1 570LDV06	eri	08-12-97 TLEV	TOYOTA: COROLLA

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EPA
1998 MODEL YEAR CERTIFICATES
Light-Duty Vehicles
with
Onboard Refueling Vapor Recovery
October 3, 1997

WVVXV2.43PA1/WVVXR0133PA1 VOLVO-LDV-98-06 TIRR 1-FED	н и	06-05-97 TLEV-CALIF	VOLVO: S70, V70. C70	S70,	V70.	020
WVVXV2,43TPF/WVVXR0133PA1 VOLVO-LDV-98-07	Ħ	07-22-97	VOLVO: S70, V70	S70,	V70	
WVVXV2.43BUP/WVVXR0133PA1 VOLVO-LDV-98-08	Ħ	1 07-22-97	VOLVO: S70, V70	570,	V70	

Appendix F

OBSERVED ORVR VEHICLES

10% ORVR Penetration

All vehicles are 1998 calendar year vehicles

2/10/98

0.47	Uaada	Accord
9. I /	nonga	Accord

- 9:35 Toyota Camry
- 10:05 Ford Escort Wagon
- 10:51 Saturn SW
- 12:20 Subaru Forester
- 13:00 Toyota Corolla
- 13:25 Chevrolet Malibu
- 15:45 Honda Accord
- 16:10 Toyota Corolla
- 17:50 Saab 900
- 17:55 Volvo \$70
- 19:15 Honda Accord
- 20:22 Pontiac Grand Prix
- 20:45 Toyota Corolla
- 23:15 Honda Accord

2/11/98

- 2:20 Buick Park Avenue
- 4:05 Ford Escort
- 7:25 Saturn SW
- 8:00 Honda Accord
- 8:35 Honda Accord
- 8:45 Buick Century
- 9:20 Mazda 626
- 9:45 Toyota Corolla
- 10: 40 Toyota Camry
- 11:15 Chevrolet Metro
- 13:30 Honda Accord
- 15:05 Oldsmobile Intrigue
- 15:20 Buick Park Avenue
- 15:50 Oldsmobile Cutlass
- 16:15 Honda Accord
- 16:25 Ford Escort

17:10 Buick Century 17:24 Hyundai Elantra 17:50 Toyota Corolla 18:25 Buick Park Avenue
19:35 Chrysler Concorde 21:20 Toyota Camry 23:15 Toyota Corolla
2/12/98
2:40 Honda Accord 5:15 Buick Century 6:30 Hyundai Elantra 7:00 Mazda 626
50% ORVR Penetration
7:40 Ford Escort Wagon 8:05 Mercury Tracer 8:30 Honda Accord 8:40 Toyota Corolla 8:55 Dodge Intrepid 9:15 Buick Regal
10:05 Buick Riviera

11:15 Honda Accord12:05 Chrysler Sebring

2:30 Chrysler Concorde3:50 Chevrolet Malibu4:10 Pontiac Grand Prix4:20 Hyundai Elantra

1:15 Ford Escort

4:45 Saab 900

7:00 Saturn SC

1:35 Honda Accord 4:00 Toyota Camry 6:25 Honda Accord

2/13/98

5:00 Toyota Corolla5:25 Subaru Legacy6:15 Subaru Impreza6:20 Chrysler Concorde

8:50 Oldsmobile Aurora 10:55 Buick Century

2:30 Chrysler Concorde3:50 Chevrolet Malibu4:10 Pontiac Grand Prix4:20 Hyundai Elantra

4:45 Saab 900

7:00 Saturn SC

2/13/98

5:00 Toyota Corolla5:25 Subaru Legacy6:15 Subaru Impreza6:20 Chrysler Concorde

8:50 Oldsmobile Aurora 10:55 Buick Century

1:35 Honda Accord

4:00 Toyota Camry 6:20 Honda Accord 6:45 Saturn SW

6:55 Chevrolet Metro

7:25 Oldsmobile Cutlass

8:00 Buick Park Avenue

8:20 Toyota Corolla 9:45 Mitsubishi Eclipse

10:05 Mercedes Benz SLK 230

11:45 Hyundai Elantra

12:30 Chevrolet Malibu

13:50 Subaru Legacy

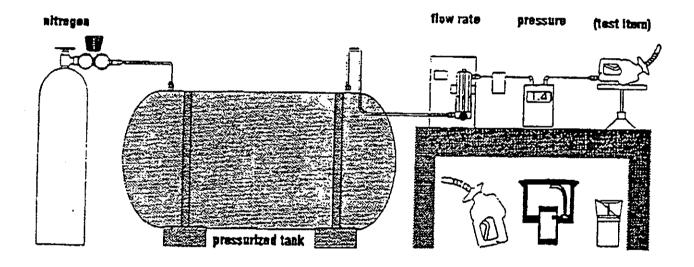
15:05 Buick Park Avenue

16:10 Ford Escort

16:35 Buick Regal

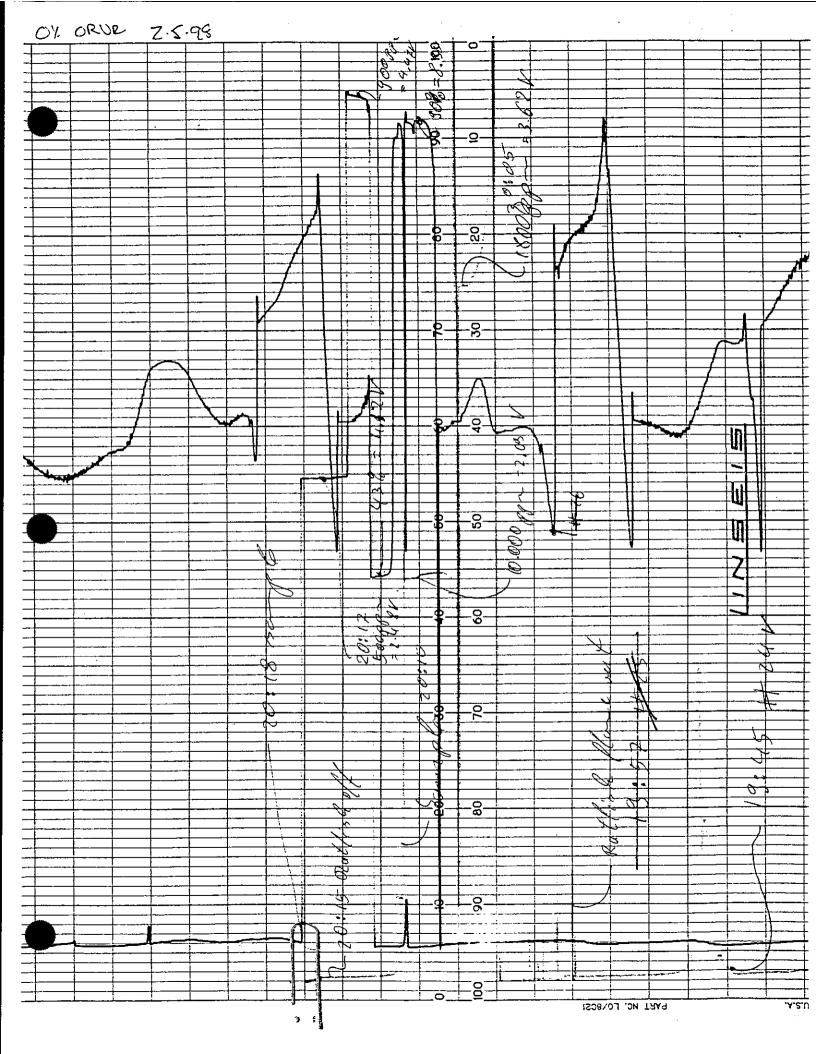
Appendix G LEAK RATE LABORATORY TESTING APPARATUS

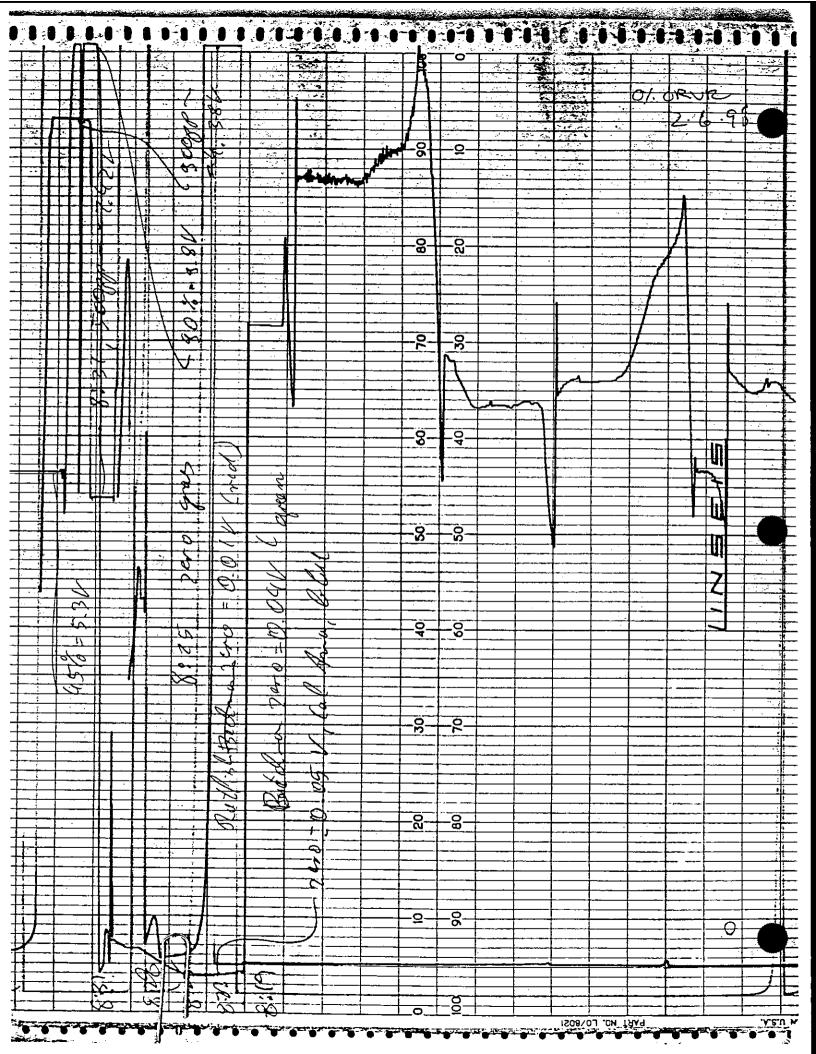
Leak Rate Laboratory Testing Apparatus

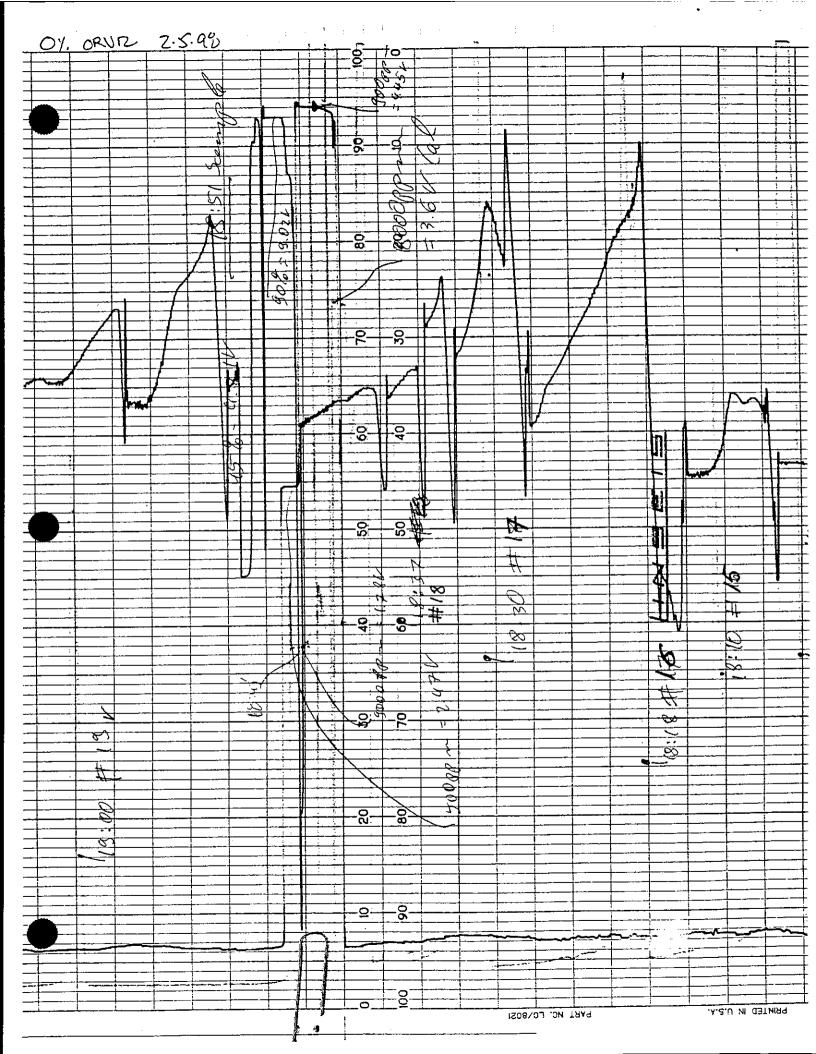


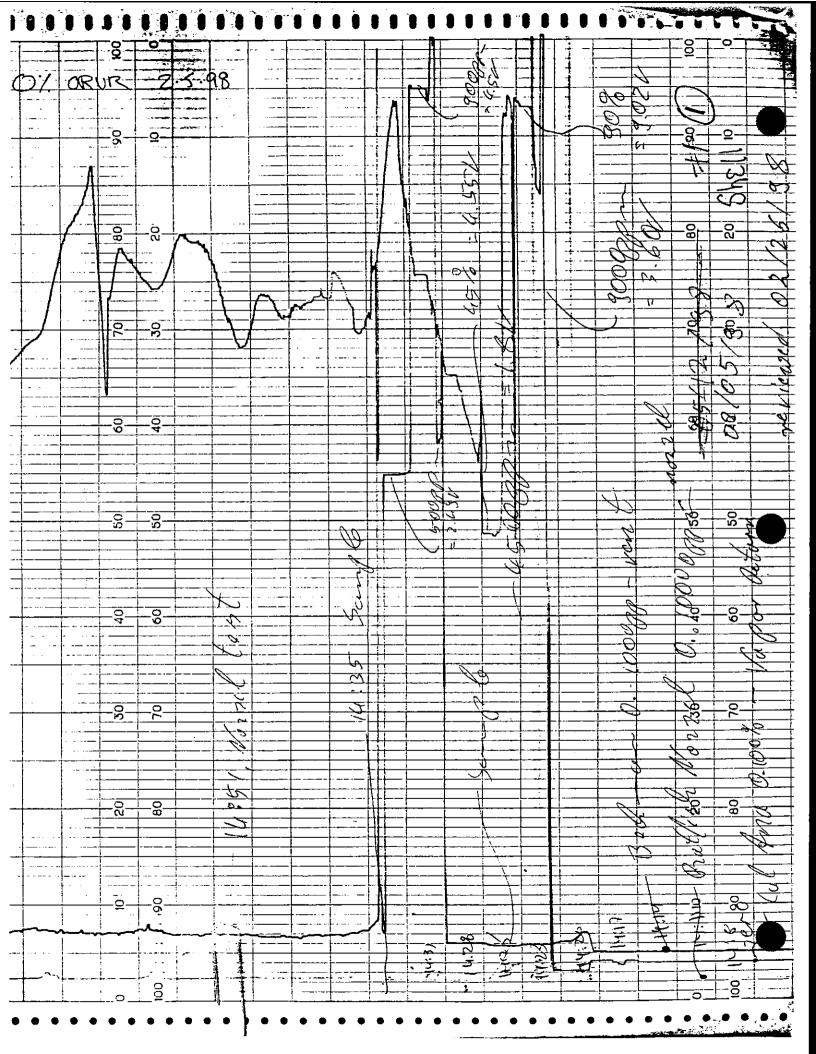
Source: Draft proposed TP-201.2B Vapor Recovery Test Procedure, Air Resources Board, January 1995

Appendix H QA/QC DATA

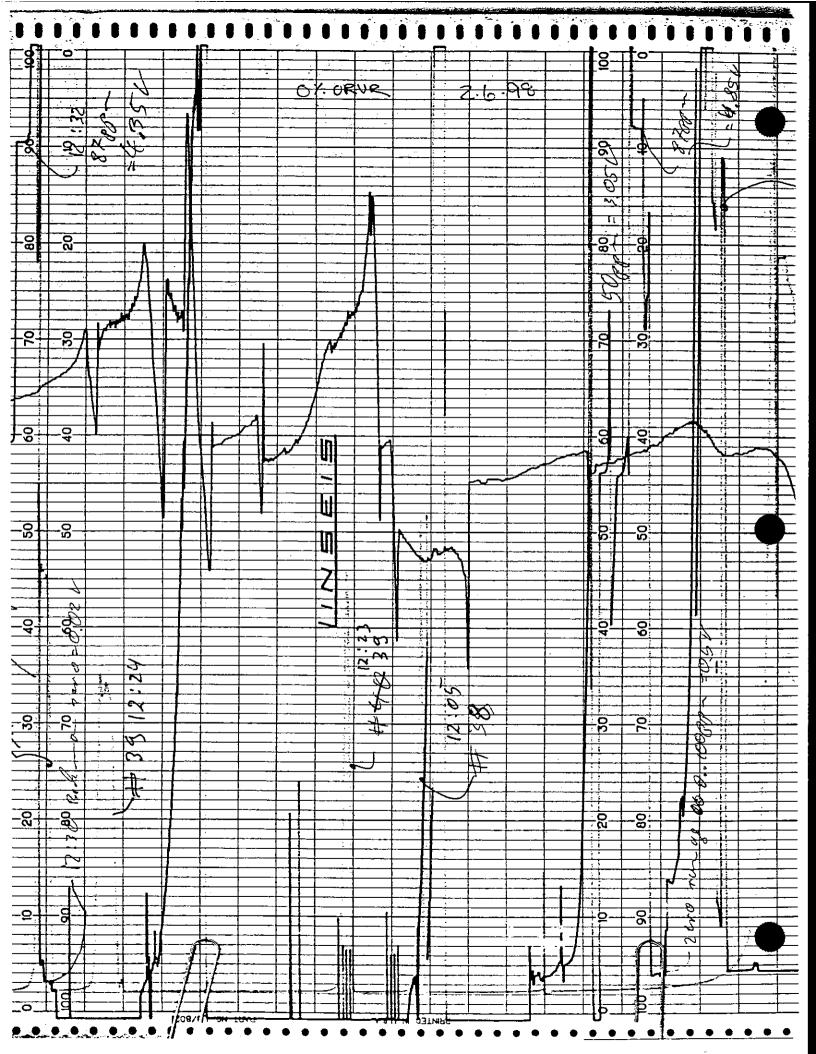


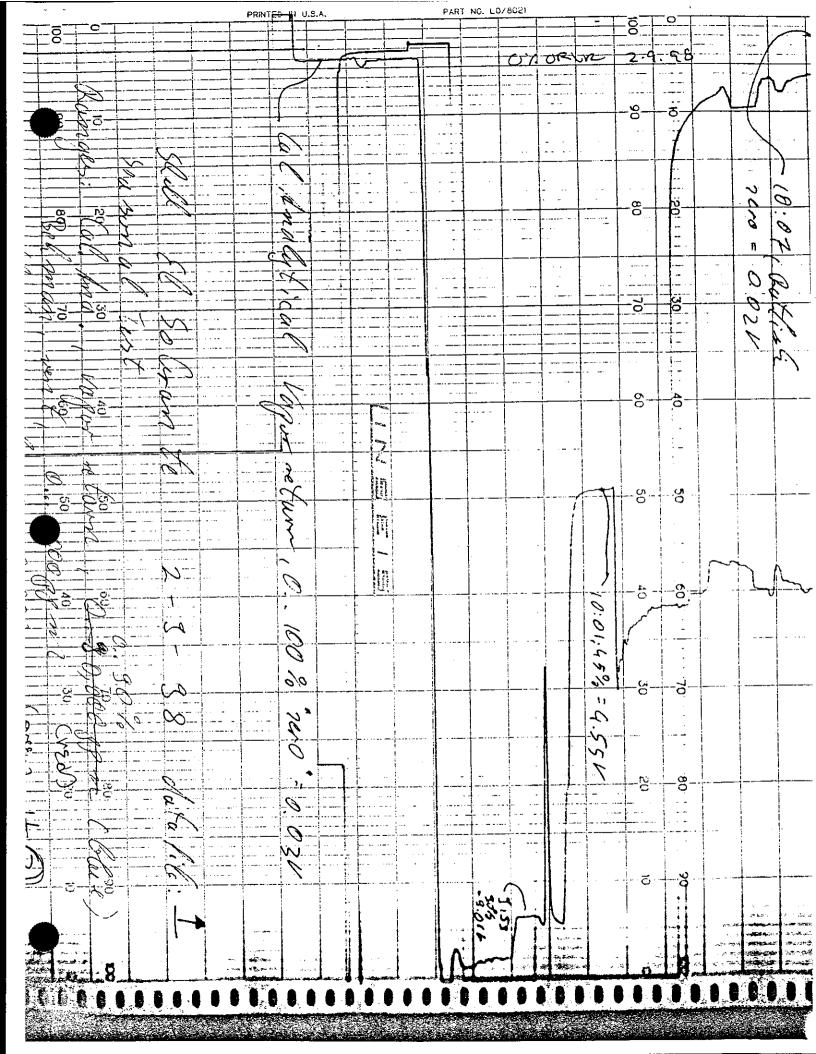


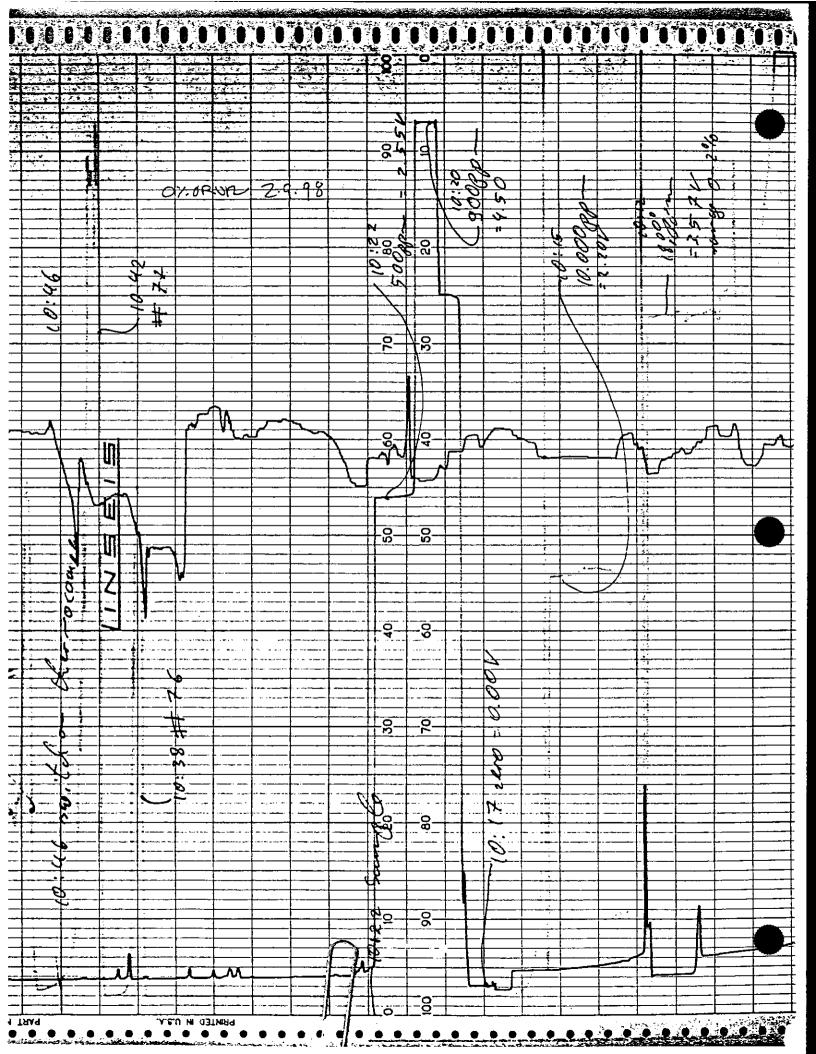


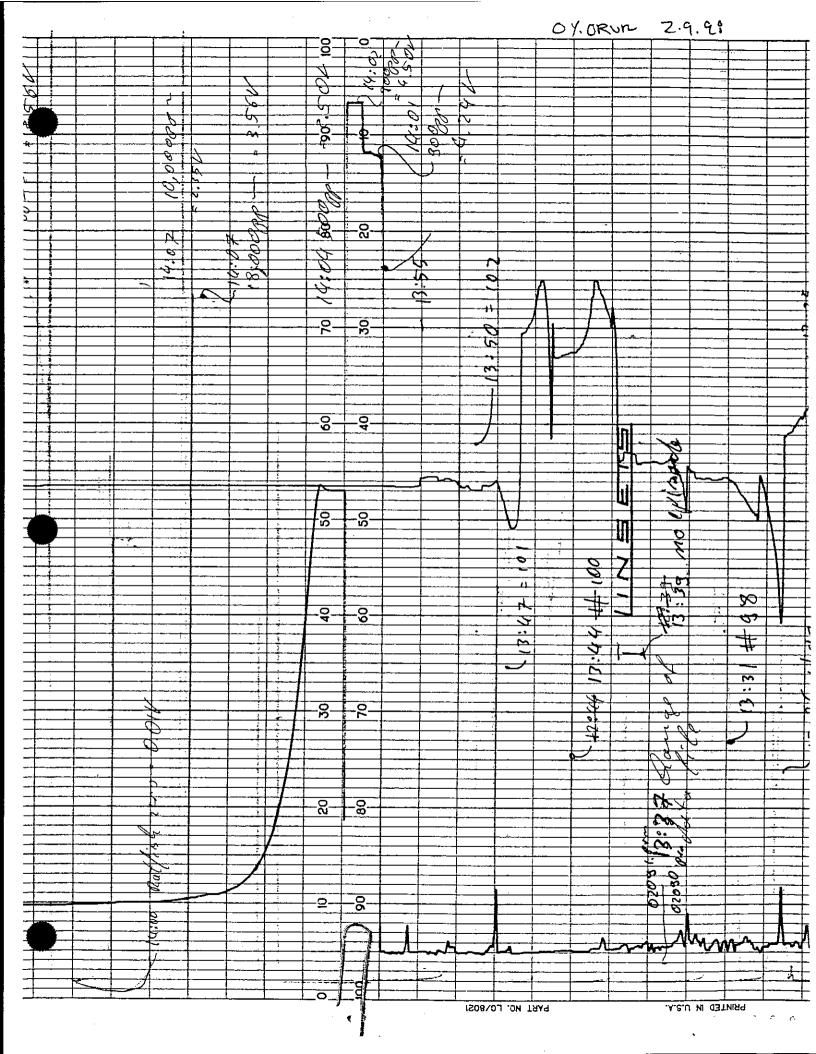


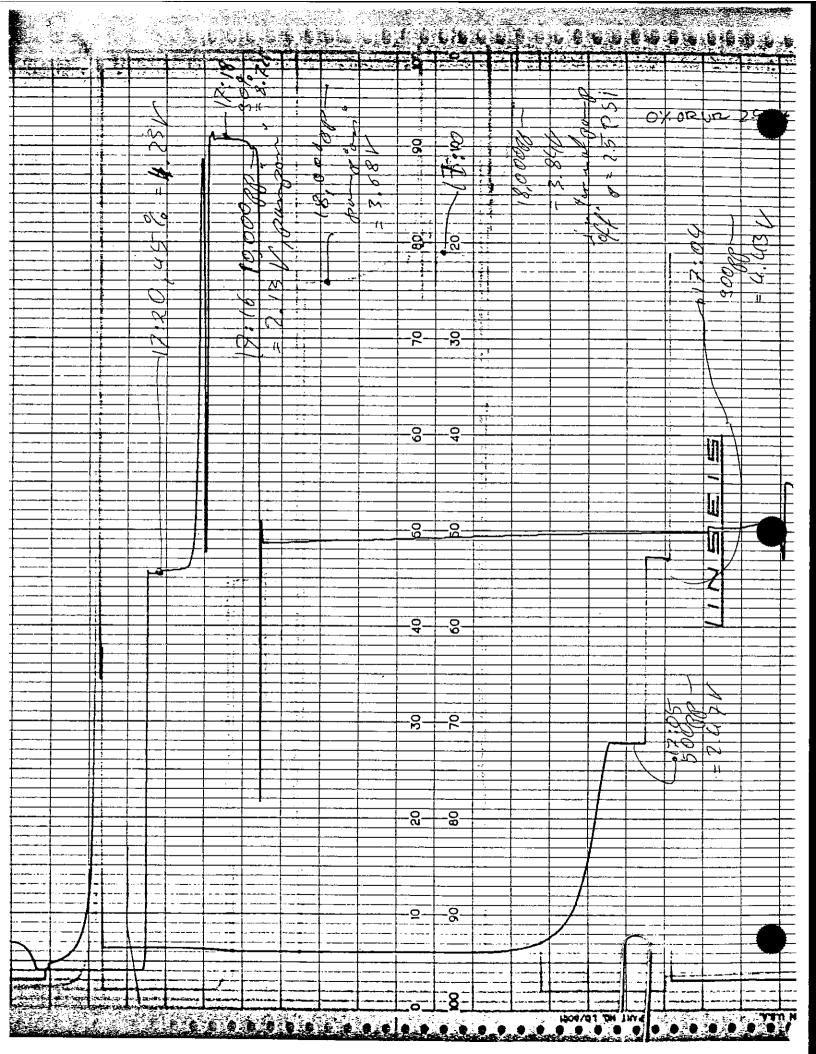
8:38	10 16 22 - 20 24 - 20
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9:50 E12 30	- 18,66 Pz-
8-3-34 G: 50: 30	
9:50 (8:43)	
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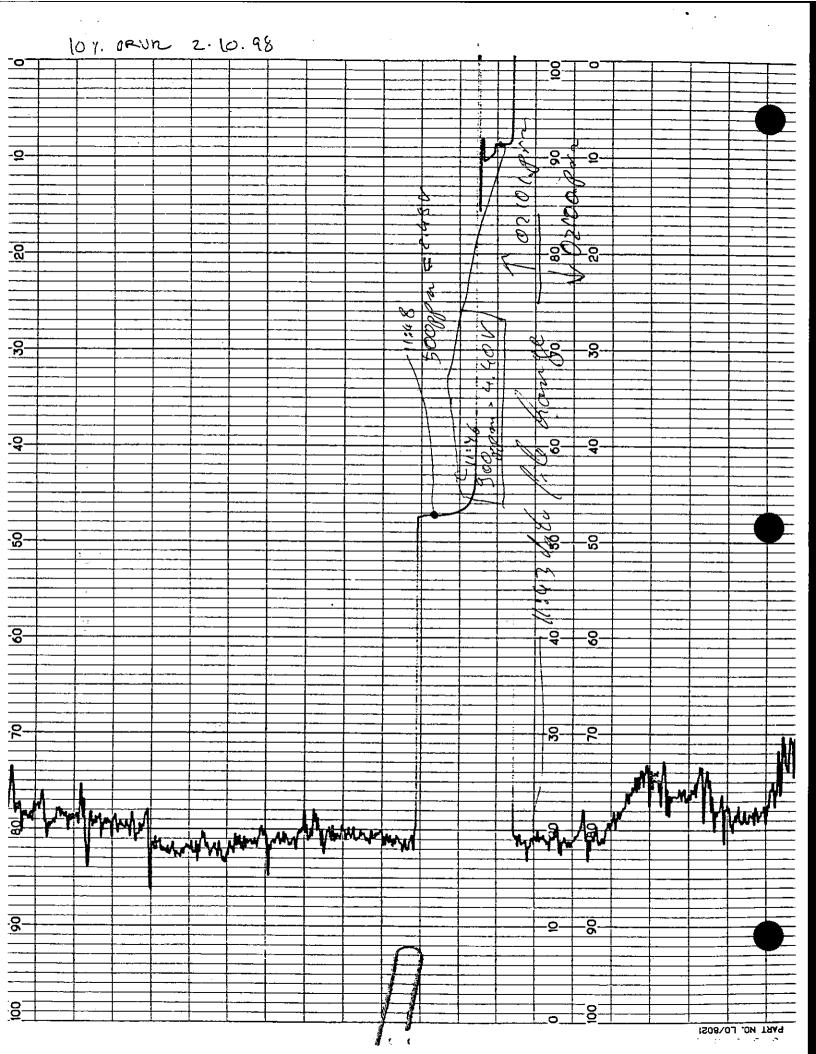


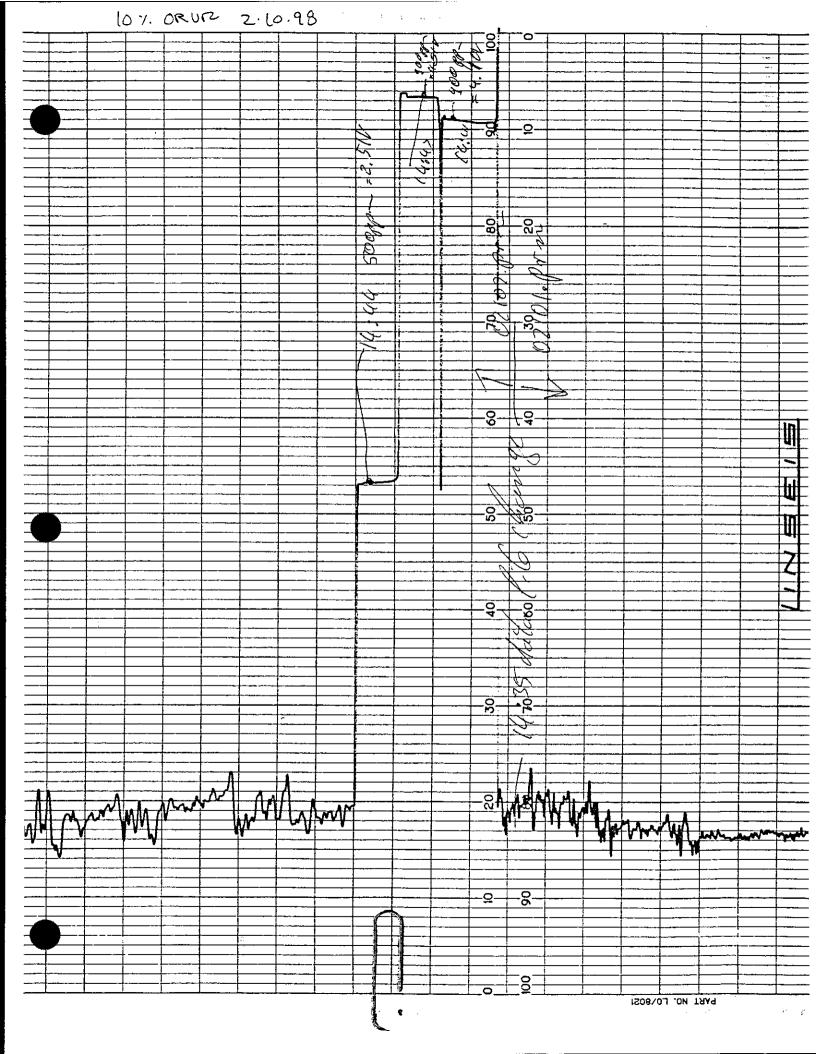


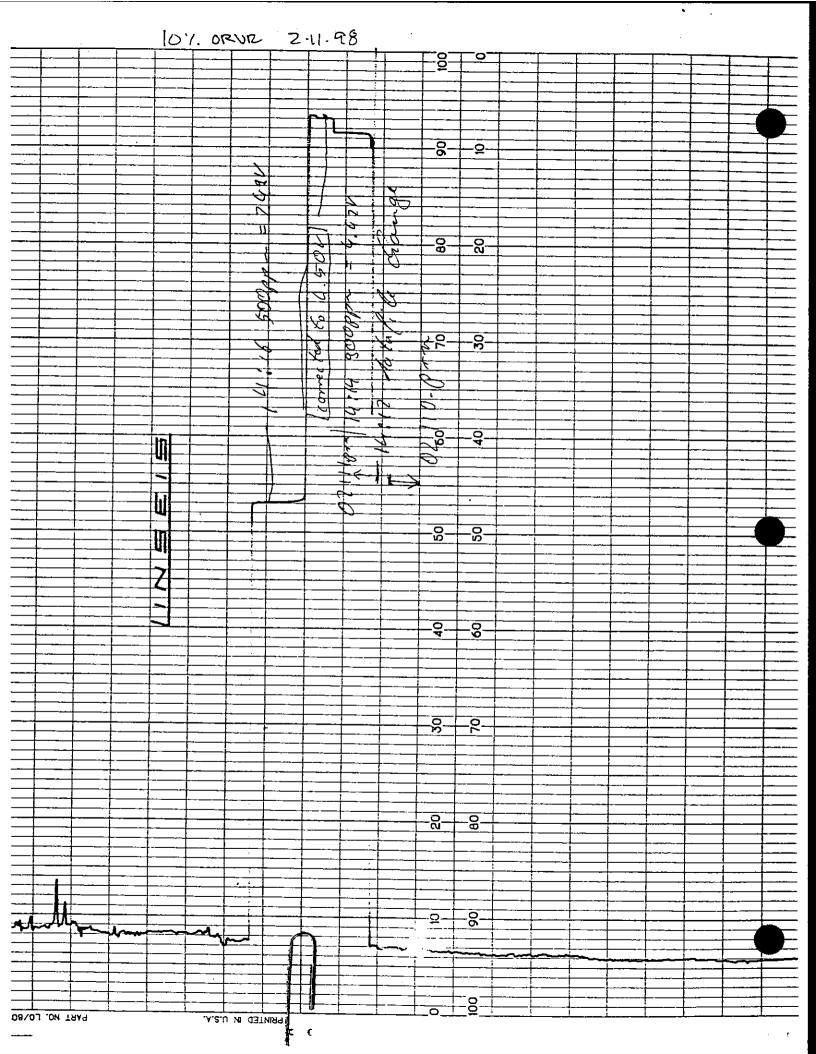


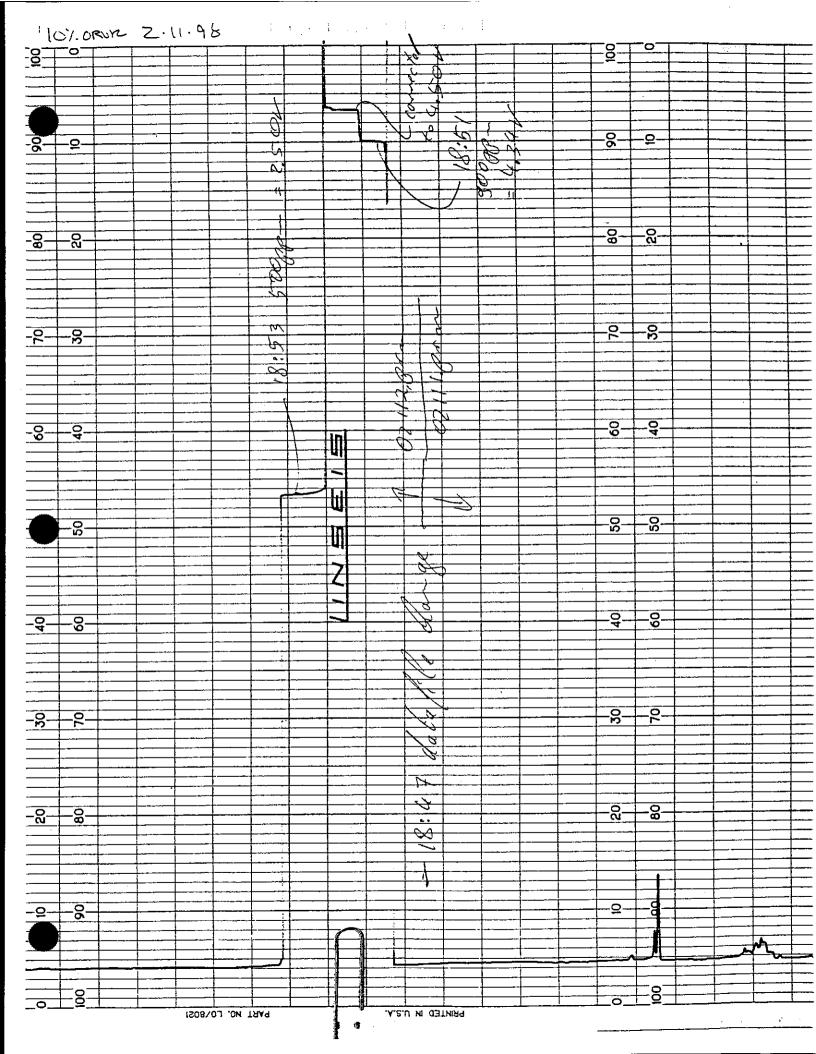


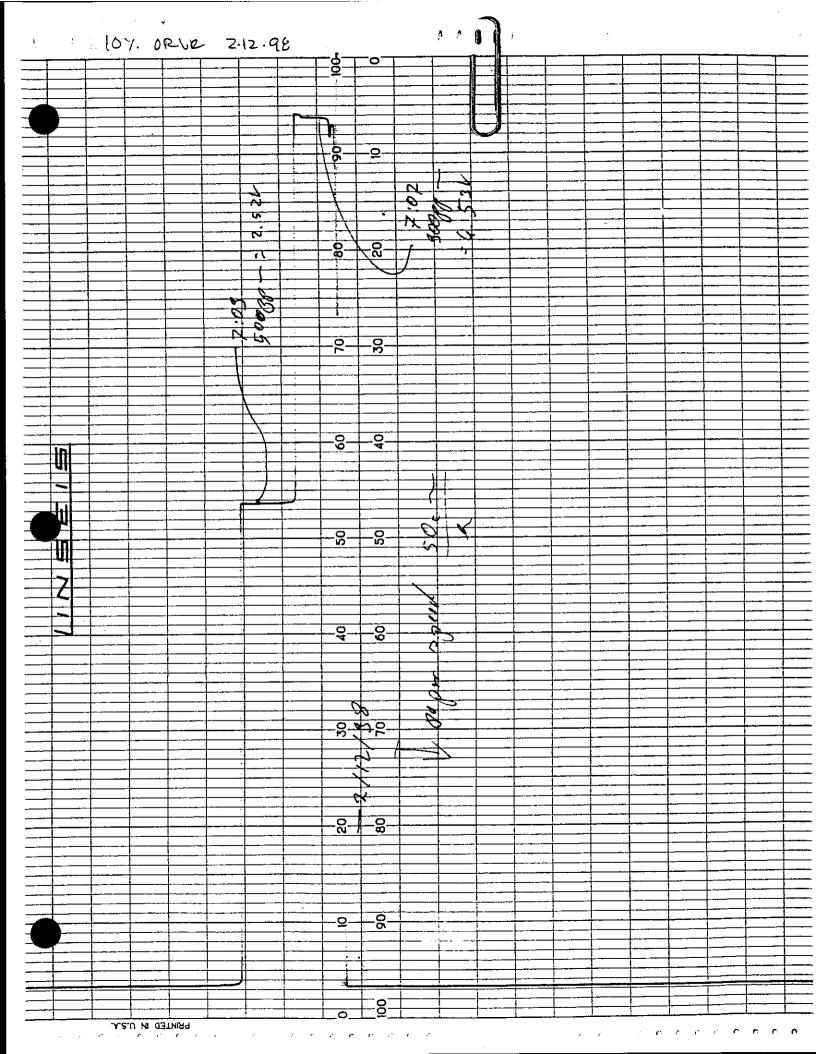


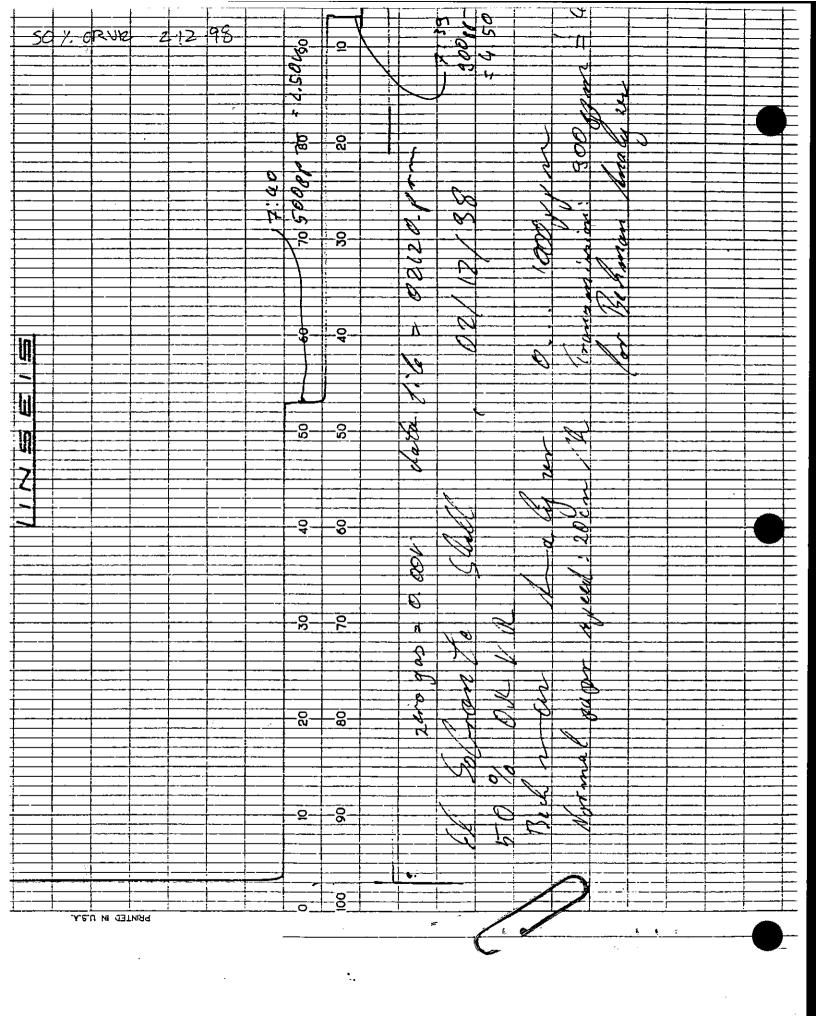












PART NO. L0/8021 -S----= -8--8--8--8--6--6-50 -5 -3--5 -3----3--8--<u>8</u>-ಕ-20 % OICALC 96.71.2

